



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Explanatory note on general fees payable to the European Medicines Agency



The fees, fee exemptions and definitions described in this explanatory note are based on *Council Regulation (EC) No 297/95 of 10.02.1995 on fees payable to the European Medicines Agency*¹ and its implementing rules (hereafter 'Fee Regulation Implementing Rules').

This explanatory note does not include any fees or charges derived from the *Regulation (EU) No 658/2014 of the European Parliament and of the Council of 15.05.2014 on fees payable to the European Medicines Agency for the conduct of pharmacovigilance activities in respect of medicinal products for human use*². For information on these fees, please see [explanatory note on pharmacovigilance fees payable to the European Medicines Agency](#).

Disclaimer:

This explanatory note is meant as a guidance note only. In case of discrepancies between the text and amounts of fees payable to the Agency quoted in the explanatory note and the provisions of the Council Regulation (EC) No 297/95, the latter document prevails.

Changes introduced in this version (1 April 2023)

- Increase in the level of fees other than administrative fees to adjust for an inflation rate of +10.4% and rounding off to the nearest EUR 100
- Increase in the level of all administrative fees to adjust for an inflation rate of +10.4% and rounding off to the nearest EUR 10
- For heading 5.6, editorial update to the reference to the Guideline on influenza vaccines – submission and procedural requirements.
- For heading 5.11, extending the scope of the fee exemptions for preparedness against biological agents that may be used as weapons of bioterrorism, from vaccines only to any medicinal products authorised under exceptional circumstances; and clarifying the conditions for applying such exemptions

¹ Official Journal L35, 15.02.1995, p. 1

² Official Journal L189, 27.06.2014, p. 112

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Medicinal products for human use

1. Medicinal products for human use

Note: This section on fees for medicinal products for human use should be read in conjunction with the section on fee exemptions (Section 5) and the annex.

1.1. Centralised procedure

1.1.1. Definitions




Pharmaceutical form: According to the 'Standard Terms' published by the Council of Europe.

Strength: See [definition](#) used in the pre-submission guidance.




Presentation: Each unit/entity of a certain strength and form of a pharmaceutical product which will be individually authorised and eventually marketed (= each individual sub-number).




1.1.2. Application for a marketing authorisation

1.1.2.1. Application for which a full dossier needs to be presented

	345 800 EURO	
Basic fee		For a single strength associated with one pharmaceutical form and one presentation.
	+ 34 800 EURO	
Additional fee		For each additional strength or pharmaceutical form including one presentation, submitted at the same time as the initial application for authorisation.
	+ 8 600 EURO	
Additional fee		For each additional presentation of the same strength and pharmaceutical form, submitted at the same time as the initial application for authorisation.

1.1.2.2. Application for which a full dossier need not be presented




	223 600 EURO	
Basic fee		For an application for a marketing authorisation pursuant to Article 10(4) of Directive 2001/83/EC. This fee is for a single strength associated with one pharmaceutical form and one presentation.
	+ 13 400 EURO	
Additional fee		For each additional strength or pharmaceutical form including one presentation submitted at the same time as the initial application for authorisation.
	+ 8 600 EURO	
Additional fee		For each additional presentation of the same strength and pharmaceutical form, submitted at the same time as the initial application for authorisation.



	134 100 EURO	
Basic fee		For applications for a marketing authorisation pursuant to Article 10(1), Article 10(3) and Article 10c of Directive 2001/83/EC. This fee is for a single strength associated with one pharmaceutical form and one presentation.
	+ 13 400 EURO	
Additional fee		For each additional strength or pharmaceutical form including one presentation submitted at the same time as the initial application for authorisation.
	+ 8 600 EURO	
Additional fee		For each additional presentation of the same strength and pharmaceutical form, submitted at the same time as the initial application for authorisation.

Note: Refer to section [A.1.1.2.](#) in the annex for examples of the determination of fees for applications for marketing authorisation.

1.1.3. Extension of a marketing authorisation

1.1.3.1. Extension of a marketing authorisation under Annex I to Commission Regulation (EC) No 1234/2008

	103 800 EURO	
Basic fee (Level I)		For each extension of a marketing authorisation within the meaning of Annex I to Commission Regulation (EC) No 1234/2008 concerning the examination of variations to the terms of a marketing authorisation for medicinal products for human use. This fee is for a single strength/potency associated with one pharmaceutical form and one presentation.
	+ 26 200 EURO	
Additional fee		For each additional strength/potency of the same pharmaceutical form of the same extension including one presentation submitted at the time of the extension application.
	+ 8 600 EURO	
Additional fee		For each additional presentation of the same strength/potency of the same pharmaceutical form of the same extension submitted at the time of the extension application.

	77 900 EURO	
Basic fee (Level II)		For all quality extensions (i.e. requiring chemical, pharmaceutical and biological documentation) for which no clinical or non-clinical data are submitted or no cross-references to previously submitted clinical or non-clinical data are made by the MAH. This fee is for a single strength/potency associated with one pharmaceutical form and one presentation. (Note: Bioequivalence data qualify as clinical data. Biowaiver dossiers are not considered as clinical data.)
	+26 200 EURO	
Additional fee		For each additional strength/potency of the same pharmaceutical form of the same extension including one presentation submitted at the time of the extension application.

**+ 8 600 EURO**

Additional fee

For each additional presentation of the same strength/potency of the same pharmaceutical form of the same extension submitted at the time of the initial extension application.

1.1.3.2. Extension of a marketing authorisation for use in the paediatric population under Article 29 of Regulation (EC) No 1901/2006

**103 800 EURO**Basic fee
(Level III)

For each extension of a marketing authorisation made under Article 29 of Regulation (EC) No 1901/2006 on medicinal products for paediatric use. This fee is for a single strength/potency associated with one pharmaceutical form and one presentation.

**+ 26 200 EURO**

Additional fee

For each additional strength/potency of the same pharmaceutical form of the same extension including one presentation submitted at the time of the extension application.

**+ 8 600 EURO**

Additional fee

For each additional presentation of the same strength/potency of the same pharmaceutical form of the same extension submitted at the time of the initial extension application.

Note: Refer to section [A.1.1.3.](#) in the annex for examples of the determination of fees for extensions of marketing authorisation.

1.1.4. Variation to a marketing authorisation

When a change affecting different strengths, pharmaceutical forms and/or presentations is introduced through a variation (e.g. to add of a new manufacturing site for active substance, to update the safety data on the product information), the corresponding fee for the variation covers all authorised strengths, pharmaceutical forms and presentations of a given medicinal product (= per main authorisation number).

The introduction of a new pack size (i.e. new Marketing Authorisation sub-number) is considered as a distinct change to the Marketing Authorisation that should be submitted as a single variation for each strength and pharmaceutical form of a given medicinal product (i.e. one variation per new Marketing Authorisation sub-number). Section [A.1.1.4.](#) in the annex includes an example of determination of fee for additional pack sizes.

The following fees apply to variations that are notified or applied for individually in a notification or application.

1.1.4.1. Type-IA variation

**3 900 EURO**

Basic fee

For a minor variation to a marketing authorisation, as defined in Article 2(2) of Commission Regulation (EC) No 1234/2008.

The fee shall be charged regardless of the outcome (positive, negative, or partial or full withdrawal).

1.1.4.2. Type-IB variation**8 600 EURO**

Basic fee

For a minor variation to a marketing authorisation, as defined in Article 2(5) of Commission Regulation (EC) No 1234/2008.

1.1.4.3. Type-II variation**103 800 EURO**Basic fee
(Level I)

For a major variation to a marketing authorisation, as defined in Article 2(3) of Commission Regulation (EC) No 1234/2008.

**77 900 EURO**Basic fee
(Level II)

For a quality variation (i.e. amendments to the chemical, pharmaceutical and biological documentation) for which no clinical or non-clinical data are submitted or no cross-references to previously submitted clinical or non-clinical data are made by the MAH. (Note: Bioequivalence data qualify as clinical data. Biowaiver dossiers are not considered as clinical data.)

**26 200 EURO**Basic fee
(Level III)

For each of the third and subsequent type II variations that are grouped in a single application made under the terms of Article 7(2) of Commission Regulation (EC) No 1234/2008.

For each of the third and subsequent type II variations to the centralised marketing authorisation or to one of the centralised marketing authorisations, where there is more than one centralised marketing authorisation, in the case of a worksharing application made under the terms of Article 20 of Commission Regulation (EC) No 1234/2008.

1.1.4.4. New indication for use in the paediatric population under Article 29 of Regulation (EC) No 1901/2006**103 800 EURO**

Basic fee

For each new indication for use in the paediatric population applied for under Article 29 of Regulation (EC) No 1901/2006 on medicinal products for paediatric use.

1.1.5. Grouping and worksharing procedures for variations**1.1.5.1. Grouping of extensions and/or variations notified or submitted under the terms of Article 7(2) of Commission Regulation (EC) No 1234/2008**

- The applicable fee specified in sections 1.1.3 and 1.1.4 shall be payable for each individual extension and/or variation to a marketing authorisation that is grouped in a single notification or a single application.
- The applicable level I and level II basic fees specified in sub-section 1.1.4.3 above are payable for the first and second type II variation respectively when both levels of fees are applicable to variations in the same grouping.
- Subsequent variations in a grouping shall be similarly charged the applicable fees as specified above.

- In the case of grouping of the same Type IA variations to the terms of several marketing authorisations owned by the same holder (as set out in Article 7(2)(a) of Commission Regulation (EC) No 1234/2008), the applicable fee shall be payable for each individual Type IA variation and for each marketing authorisation in the grouping.
- The same marketing authorisation holder also means several marketing authorisation holders that are linked through a parent company. The fee for the grouping shall be payable by the marketing authorisation holder applying for the grouping procedure.
- Where any extensions/variations included in a grouping are found not to be valid and the remainder are validated positively, the applicable fees as specified above shall be payable for each of the positively validated extensions/variations.

1.1.5.2. Worksharing procedure under the terms of Article 20 of Commission Regulation (EC) No 1234/2008


- The applicable fee specified in section 1.1.4 is payable for each individual variation to one of the centralised marketing authorisations owned by the same holder, where more than one centralised marketing authorisation is included in the worksharing application, or to the single centralised marketing authorisation included in the worksharing application, as applicable.
- The applicable level I and level II basic fees specified in 1.1.4.3 above are payable for the first and second type II variation respectively when both levels of fees are applicable to variations in the same worksharing procedure.
- The administrative fee shown in the table below is additionally payable for each individual variation to each of the other centralised marketing authorisation(s) owned by the same holder included in the same worksharing application, if applicable.
- No fee is payable to the Agency for any national marketing authorisations included in the same worksharing application.
- The same marketing authorisation holder also means several marketing authorisation holders that are linked through a parent company. The fee for the worksharing procedure shall be payable by the marketing authorisation holder applying for the procedure.

Variation type	Worksharing fees for one centralised marketing authorisation	Worksharing administrative fees for second and subsequent marketing authorisations	
		Excluding applications on usage patent grounds	Applications on usage patent grounds ³
Type II (Level I)	103 800 EURO	8 660 EURO	4 980 EURO
Type II (Level II)	77 900 EURO		
Type II (Level III)	26 200 EURO		
Type IB	8 600 EURO	1 450 EURO	
Type IA	3 900 EURO	720 EURO	

³ These fees are applicable for as long as the concerned marketing authorisation is affected by usage patent(s) pertaining to indication(s) and/or dosage form(s). Refer to section 4.3 for further details on fee exemptions.




Where any variations included in a worksharing procedure are found not to be valid and the remainder are validated positively, the applicable fees as specified above shall be payable for each of the positively validated variations.

1.1.6. Renewal of a marketing authorisation

	17 000 EURO
Basic fee	For each strength associated with a pharmaceutical form for which renewal is sought.


Note: Refer to section [A.1.1.6.](#) in the annex for examples of the determination of fees for renewals of marketing authorisation.

1.1.7. Inspection


	26 200 EURO
Basic fee (Level I)	For each inspection inside or outside the European Union; for inspections outside the European Union, travel expenses shall be charged extra on the basis of actual cost.
	13 000 EURO
Basic fee (Level II)	For each consecutive distinct plasma master file (PMF) inspection performed in conjunction with an inspection that attracts the level I fee, provided that such consecutive inspection concerns the same PMF application, the same inspection team and is conducted in the same PMF inspection tour.
	13 000 EURO
Basic fee (Level III)	For each inspection inside or outside the European Union cancelled due to the withdrawal of the application or intended application; or changes to the manufacturing arrangements made by the manufacturer or changes made by the applicant/prospective applicant/MAH that necessitate a cancellation of the inspection before the inspection is carried out.

Note: Refer to section [A.1.1.7.1.](#) in the annex for examples of the determination of fees for good manufacturing practice (GMP) inspections and to section [A.1.1.7.2.](#) in the annex for examples of the determination of fees for good clinical practise (GCP).

1.1.8. Transfer of a marketing authorisation

	8 600 EURO
Basic fee	For all authorised presentations of the medicinal product.

1.1.9. Maintenance of a marketing authorisation – Annual fee

	123 900 EURO
Basic fee (Level I)	For each marketing authorisation of a medicinal product. This fee covers all authorised presentations of the medicinal product.



62 000 EURO

Basic fee
(Level II)

For each marketing authorisation of a biosimilar medicinal product (Article 10(4) of Directive 2001/83/EC). This fee covers all authorised presentations of the medicinal product.



31 000 EURO

Basic fee
(Level III)

For each marketing authorisation of a generic, hybrid or informed consent medicinal product (Articles 10(1), 10(3) and 10c of Directive 2001/83/EC). This fee covers all authorised presentations of the medicinal product.

1.1.10. Referral



86 000 EURO

Basic fee

For procedures laid down in Article 30(1) and 31 of Directive 2001/83/EC that are initiated by the applicant of a marketing authorisation or the holder of an existing marketing authorisation.

Note: Where more than one applicant for marketing authorisation or holder of existing marketing authorisations are concerned by the procedures referred to in the first subparagraph, the applicants or holders may be grouped for the purpose of the payment of one single referral fee. If, however, the same procedure concerns more than 10 different applicants or holders, the fee will be charged by the application of the above-mentioned referral fee.

The fee for procedures, which are initiated according to the second subparagraph of Article 31(1) of Directive 2001/83/EC, will be charged according to Article 6 of Regulation (EU) 658/2014 of 15 May 2014 (pharmacovigilance fee regulation).

1.2. Scientific advice

1.2.1. Definitions

Quality development: Chemical, pharmaceutical and biological testing.

Safety development: Toxicological and pharmacological tests.

Clinical development: Studies in human subjects, whether patients or non-patient volunteers, including clinical pharmacological trials designed to determine the efficacy and safety of the product; may also include guidance to demonstrate significant benefit over authorised medicines in the context of a designated orphan medicinal product.

Qualification advice: Advice on the acceptability of a specific use of the proposed method (e.g. use of a biomarker) in a research and development (R&D) context (non-clinical or clinical studies), based on the assessment of submitted data.

Initial request: First request for scientific advice or protocol assistance introduced in relation to the submission of an application for marketing authorisation or a variation, whatever the authorisation phase (pre- or post-authorisation).

Follow-up to initial request: Any subsequent request falling within the same therapeutic indication and initial area(s) as the initial request, (area meaning quality, preclinical and/or clinical development, including pharmacovigilance/risk management aspects).

1.2.2. Initial request for scientific advice



51 800 EURO

Basic fee
(Level I)

For **initial** requests on:

- quality development or
- safety development, or
- bioequivalence studies for generic medicinal products.



77 900 EURO

Basic fee
(Level II)

For **initial** requests on:

- clinical development, or
- quality and safety development, or
- quality and bioequivalence studies for generic medicinal products.



103 800 EURO

Basic fee
(Level III)

For **initial** requests for scientific advice on:

- Quality, and safety and clinical development, or
- quality and clinical development, or
- safety and clinical development, or
- qualification advice.

1.2.3. Follow-up request for scientific advice



26 200 EURO

Basic fee
(Level I)

For **follow-up** to the initial request on:

- quality development, or
- safety development, or
- bioequivalence studies for generic medicinal products.



39 100 EURO

Basic fee
(Level II)

For **follow-up** to the initial request on:

- clinical development, or
- quality and safety development, or
- quality and bioequivalence studies for generic medicinal products.



51 800 EURO

Basic fee
(Level III)

For **follow-up** to the initial request on:

- quality and safety and clinical development, or
- quality and clinical development, or
- safety and clinical development, or
- qualification advice.

1.3. Scientific services

A scientific service fee shall apply where an application is made for any scientific advice or opinion by a scientific Committee, which is not covered by Articles 3 to 7 or by Article 8(1).

1.3.1. Scientific opinions pursuant to Article 58 of Regulation (EC) No 726/2004

The basic and additional fees specified in section 1.1.2 apply by analogy for a scientific opinion for the evaluation of medicinal products for human use intended exclusively for markets outside the European Union. Likewise, the inspection fees in section 1.1.7 apply by analogy to any inspection undertaken for the purpose of assessment prior to an opinion.

Fees for post-opinion services and annual fees are charged according to the corresponding fees for centrally authorised products. Therefore, the fees specified in sections 1.1.3, 1.1.4, 1.1.5, 1.1.7 and 1.1.9 apply by analogy.

The fee incentives for micro, small and medium-sized enterprises apply to scientific services described in section 5.1.2. However, fee deferrals and conditional fee exemptions do not apply to services in relation to scientific opinions pursuant to Article 58 of Regulation (EC) No 726/2004.

1.3.2. Compassionate use



173 000 EURO

Basic fee

For any opinion on medicinal products for compassionate use.

This fee will be deducted from the fee charged for a subsequent application for marketing authorisation by the same applicant.

1.3.3. Herbal medicinal products



26 200 EURO

Basic fee
(Level I)

For requests for scientific support and advice by the Committee on Herbal Medicinal Products (HMPC) on multiple areas related to traditional herbal medicinal products.



17 000 EURO

Basic fee
(Level II)

For requests for scientific support and advice by the HMPC on single areas related to traditional herbal medicinal products, for example:

- quality, or
- safety, or
- long-standing use.

1.3.4. Certification of quality and non-clinical data relating to advanced therapy medicinal products developed by micro, small and medium-sized enterprises



77 900 EURO

Basic fee

Evaluation of an application relating to quality and non-clinical data.

(Level I)

**51 800 EURO**Basic fee
(Level II)

Evaluation of an application relating to quality data.

1.3.5. Rolling Review

**173 000 EURO**

Basic fee

For any application for a 'Rolling Review' Assessment for potential marketing authorisations in a human pandemic situation.

This fee will be deducted from the fee charged for a subsequent application for marketing authorisation by the same applicant. In case of multiple Rolling Review cycles this fee will only be charged once.

1.4. Consultation on medical devices

For the establishment of fee fees applying to consultation on medical device, the following definitions are applied:

- *Initial request:* first request for consultation on a medical device.
- *Follow-up to initial request:* any subsequent request for a consultation on a device that previously underwent an initial consultation with the EMA.

The fee incentives for micro, small and medium-sized enterprises apply to scientific services described in section [5.1.2.](#)

The fees payable for consultations on medical devices shall be charged to the medical device manufacturer that, according to the application form submitted to the Agency, requested the assessment of conformity of the medical device by the notified body on the basis of which the consultation is applied for.

1.4.1. Ancillary substances including blood derivatives incorporated in medical devices

The determination of the fee shall be guided by the following principles:

- Where a device incorporates two or more ancillary substances/derivatives, the fee relates to one of the substances/derivatives only – the one that commands the highest fee.
- One application may include a range of strengths or concentrations of the ancillary substance/derivative and/or a range of similar devices (e.g. a range of catheters made of the same material) incorporating the same substance/derivative from the same manufacturer.

1.4.1.1. Initial request

**103 800 EURO**Basic fee
(Level I)

For consultation on an ancillary medicinal substance or blood derivative new to the centralised procedure. This fee applies where the substance/derivative from the specified manufacturer has not been evaluated by the Agency in connection with a previous marketing authorisation and/or through a previous successful

notified body consultation.



77 900 EURO

Basic fee
(Level II)

For consultation on a known ancillary blood derivative from a known source. This fee applies where the blood derivative from the specified manufacturer has been evaluated by the Agency in connection with a previous marketing authorisation and/or through a previous successful notified body consultation.



51 800 EURO

Basic fee
(Level III)

For consultation on a known ancillary medicinal substance from a known source. This fee applies where the substance from the specified manufacturer has been evaluated by the Agency in connection with a previous marketing authorisation and/or through a previous successful notified body consultation.

1.4.1.2. Follow-up to the initial request



51 800 EURO

Basic fee

For consultation on an amendment to the documentation on an ancillary medicinal substance or blood derivative already evaluated by the Agency (The amendments should be classified by analogy to Annex I of Commission Regulation (EC) No 1234/2008).



51 800 EURO

Basic fee

For consultation on a major amendment to the documentation on an ancillary medicinal substance or blood derivative already evaluated by the Agency (The amendment should be equivalent to a Type II variation as classified by analogy to Commission Regulation (EC) No 1234/2008).



51 800 EURO

Basic fee

For consultation on two or more amendments to the documentation on an ancillary medicinal substance or blood derivative already evaluated by the Agency that are grouped in a single application by analogy to the terms of Articles 7(2)(b) or 7(2)(c) of Commission Regulation (EC) No 1234/2008 where at least one of the amendments is classified by analogy to Annex I of Commission Regulation (EC) No 1234/2008 or as a major amendment.



8 600 EURO

Basic fee

For consultation on a minor amendment to the documentation on an ancillary medicinal substance or blood derivative already evaluated by the Agency. (The amendment should be equivalent to a Type IB variation as classified by analogy to Commission Regulation (EC) No 1234/2008).



3 900 EURO

Basic fee

For consultation on a minor amendment to the documentation on an ancillary medicinal substance or blood derivative already evaluated by the Agency. (The amendment should be equivalent to a Type IA variation as classified by analogy to Commission Regulation (EC) No 1234/2008).

1.4.2. Medical devices composed of substances or combinations of substances that are systemically absorbed to achieve their intended purpose

1.4.2.1. Initial request

The determination of the fee shall be guided by the following principles:

- Where a device is composed of two or more substances the consultation commands one fee only.
- One application may include a range of presentations of the substance(s) from the same manufacturer.



77 900 EURO

Basic fee

For consultation on the compliance of medical devices composed of substance(s) or combination of substances that are systemically absorbed to achieve their intended purpose with the relevant requirements of Annex I to Directive 2001/83/EC.

1.4.3. Companion diagnostics

1.4.3.1. Initial request

The determination of the fee shall be guided by the following principles:

- The consultation will be charged for suitability of the device to medicinal product(s) containing the same active substance(s) or the same combination of active substances⁴
- Applications for consultation submitted simultaneously for suitability of the device in relation to more than one active substance(s) or combination of substances will incur an added charge per additional active substance or combination of substances up to a maximum fee.



51 800 EURO

Basic fee

For consultation on the suitability of a companion diagnostic in relation to a concerned medicinal product. This fee applies in case a consultation is requested by a notified body to support a new conformity assessment.

The consultation will be charged at fee of 51 800 EURO for suitability of the device to medicinal product(s) containing the same active substance(s) or the same combination of active substances.



+ 8 600 EURO

Additional fee

For applications submitted simultaneously for suitability of the device in relation to each additional active substance or combination of substances up to a maximum total fee of 103 800 EURO.

⁴ In the context of the consultation for companion diagnostics, the different salts, esters, ethers, isomers, mixtures of isomers, complexes or derivatives of an active substance shall be considered to be the same active substance, unless they differ significantly in properties with regard to safety and/or efficacy.

1.4.3.2. Follow-up to the initial request



26 200 EURO

Basic fee

For consultation on an amendment to the documentation of a companion diagnostic already evaluated by the Agency in connection with a previous successful notified body consultation. In this case a further consultation is requested by a notified body after a first consultation, in response to changes to a companion diagnostic that affect the performance and/or the intended use and/or the suitability of the device in relation to the medicinal product(s) concerned, which results in a supplement to the EU technical documentation assessment certificate or EU type-examination certificate.

1.5. Certification of compliance with European Union legislation for plasma master files (PMF) and vaccine antigen master files (VAMF)

1.5.1. Application for PMF certification (initial certification)

1.5.1.1. Not submitted simultaneously with a new application under the centralised procedure



86 000 EURO

Basic fee
(Level I)

For the review of the PMF and its initial certification where the data contained in the PMF have not been previously evaluated within the centralised procedure.



77 900 EURO

Basic fee
(Level II)

For the review of the PMF and its initial certification where the PMF applicant has included change(s) to the data previously evaluated within the centralised procedure and which are now part of the PMF application.



26 200 EURO

Basic fee
(Level III)

For the review of the PMF and its initial certification where the data contained in the PMF have been previously evaluated under the centralised procedure and no changes have been included.

1.5.1.2. Submitted simultaneously with a new application under the centralised procedure



8 600 EURO

Basic fee

For the review of the PMF and its certification when it is submitted in parallel and within the submission of a new application within the centralised procedure. The PMF documentation will be evaluated by the Agency simultaneously with a centralised marketing authorisation application.

1.5.2. Variation to a certified PMF



77 900 EURO

Basic fee

For the review and certification of a major variation to the PMF (in accordance with Commission Regulation (EC) No 1234/2008).



77 900 EURO

Basic fee

For the review and certification of two or more variations that are grouped in a single application made under the terms of Articles 7(2)(b) or 7(2)(c) of

Commission Regulation (EC) No 1234/2008 where at least one of the variations is a major variation.



8 600 EURO

Basic fee

For the review and certification of a minor variation of type IB to the PMF (in accordance with Commission Regulation (EC) No 1234/2008).



3 900 EURO

Basic fee

For the review and certification of a minor variation of type IA to the PMF (in accordance with Commission Regulation (EC) No 1234/2008).

1.5.3. Annual re-certification of PMF



77 900 EURO

Basic fee

For the review and annual re-certification of the PMF under this scheme where one or more variations are included in the submitted documentation and at least one of the variations is a major variation.



17 000 EURO

Basic fee

For the review and annual re-certification of the PMF under this scheme where no major variations are included in the submitted documentation.

This fee is increased by the applicable fee for each minor variation of type IA or type IB included in the submitted documentation, up to a maximum total fee of 77 900 EURO.

1.5.4. Application for a VAMF certification (initial certification)

1.5.4.1. Not submitted simultaneously with a new application under the centralised procedure



86 000 EURO

Basic fee

For the review of the VAMF and its certification where the data contained in the vaccine antigen master file have not been previously evaluated within the centralised procedure. In the case of a group of antigens aimed at preventing a single infectious disease, e.g. inactivated polio serotypes 1, 2 and 3, the VAMF application for one antigen will be charged at the above fee.



+ 8 600 EURO

Additional fee

For each VAMF application submitted simultaneously for antigens from the same group, up to a maximum total fee of 103 800 EURO.





77 900 EURO


Basic fee

For the review of the VAMF and its certification where the initial data have been previously evaluated within the centralised procedure but where the VAMF applicant has included changes or harmonisation as part of the VAMF certification scheme.


In the case of a group of antigens aimed at preventing a single infectious disease e.g. inactivated polio serotypes 1, 2 and 3, the VAMF application for one antigen will be charged at the above fee.


 **+ 8 600 EURO**
 Additional fee For each VAMF application submitted simultaneously for antigens from the same group, up to a maximum total fee of 103 800 EURO.

 **26 200 EURO**
 Basic fee For the review of the VAMF and its initial certification where the data contained in the vaccine antigen master file have been previously evaluated under the centralised procedure and where no changes or harmonisation have been included.
 In the case of a group of antigens aimed at preventing a single infectious disease e.g. Inactivated Polio serotypes 1, 2 and 3, the VAMF application for one antigen will be charged at the above fee.


 **+ 8 600 EURO**
 Additional fee For each VAMF application submitted simultaneously for antigens from the same group, up to a maximum total fee of 103 800 EURO.


1.5.4.2. Submitted simultaneously with a new application under the centralised procedure


 **8 600 EURO**
 Basic fee For the review of the VAMF and its certification when it is submitted in parallel and within the submission of a new application containing the named antigen within the centralised procedure. In the case of a group of antigens aimed at preventing a single infectious disease e.g. inactivated polio serotypes 1, 2 and 3, the VAMF application for one antigen will be charged at the above fee.

 **+ 8 600 EURO**
 Additional fee For each VAMF application submitted simultaneously for antigens from the same group, up to a maximum total fee of 103 800 EURO.

1.5.5. Variation to a certified VAMF

 **77 900 EURO**
 Basic fee For the review and certification of a major variation to the VAMF (in accordance with Commission Regulation (EC) No 1234/2008).
 In the case of a group of antigens aimed at preventing a single infectious disease e.g. inactivated polio serotypes 1, 2 and 3, the VAMF application for one antigen will be charged at the above fee.

 **+ 8 600 EURO**
 Additional fee For each VAMF application submitted simultaneously for antigens from the same group, up to a maximum total fee of 103 800 EURO.

 **77 900 EURO**
 Basic fee For the review and certification of two or more variations that are grouped in a single application made under the terms of Articles 7(2)(b) or 7(2)(c) of Commission Regulation (EC) No 1234/2008 where at least one of the variations is a major variation.

In the case of a group of antigens aimed at preventing a single infectious disease e.g. inactivated polio serotypes 1, 2 and 3, the grouped variations application for one antigen will be charged at the above fee.



+ 8 600 EURO

Additional fee

For each VAMF grouping application submitted simultaneously for antigens from the same group, up to a maximum total fee of 103 800 EURO.



8 600 EURO

Basic fee

For the review and certification of a minor variation of type IB to the VAMF (in accordance with Commission Regulation (EC) No 1234/2008).

In the case of a group of antigens aimed at preventing a single infectious disease e.g. inactivated polio serotypes 1, 2 and 3, the VAMF application for one antigen will be charged at the above fee.



+ 8 600 EURO

Additional fee

For each VAMF applications submitted simultaneously for antigens from the same group, up to a maximum total fee of 51 800 EURO.



3 900 EURO

Basic fee

For the review and certification of a minor variation of type IA to the VAMF (in accordance with Commission Regulation (EC) No 1234/2008).

In the case of a group of antigens aimed at preventing a single infectious disease e.g. inactivated polio serotypes 1, 2 and 3, the VAMF application for one antigen will be charged at the above fee.



+ 3 900 EURO

Additional fee

For each VAMF application submitted simultaneously for antigens from the same group, up to a maximum total fee of 26 200 EURO.



Veterinary medicinal products

2. Veterinary medicinal products

Note: This section on fees for veterinary medicinal products should be read in conjunction with the section on fee exemptions (Section 5) and the annex.

2.1. Centralised procedure

2.1.1. Definitions



Pharmaceutical form: According to the 'Standard Terms' published by the Council of Europe.

Strength: See [definition](#) used in the pre-submission guidance.

Presentation: Each unit/entity of a certain strength and form of a pharmaceutical product which will be individually authorised and eventually marketed (= each individual sub-number).




2.1.2. Application for a marketing authorisation


2.1.2.1. Full fee

	173 000 EURO
Basic fee	For a single strength associated with one pharmaceutical form and one presentation.
	+ 17 000 EURO
Additional fee	For each additional strength or pharmaceutical form including one presentation, submitted at the same time as the initial application for authorisation.
	+ 8 600 EURO
	For each additional presentation of the same strength and pharmaceutical form, submitted at the same time as the initial application for authorisation.

Number of target species applied for does not impact on the fee.




2.1.2.2. Full fee – Immunologicals




	86 000 EURO
Basic fee	For a single strength associated with one pharmaceutical form and one presentation.
	+ 8 600 EURO
Additional fee (I)	For each additional strength or pharmaceutical form including one presentation submitted at the same time as the initial application for authorisation.
	+ 8 600 EURO
Additional fee (I)	For each additional presentation of the same strength and pharmaceutical form, submitted at the same time as the initial application for authorisation.

 Additional fee (II)	<p>+ 8 600 EURO</p> <p>For each multi-strain additional presentation of the same application submitted at the time of the initial application up to a maximum total fee of 173 000 EURO.</p> <p>In this context, each combination of strain identified in the application represents a presentation.</p>
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The number of target species applied for does not impact on the fee.




2.1.2.3. Reduced fee

 Basic fee	<p>86 000 EURO</p> <p>For applications for marketing authorisation pursuant to Article 18, 19 (except for the veterinary medicinal products covered in the table below) 21 and 23 of Regulation (EU) 2019/6. This fee is for a single strength associated with one pharmaceutical form and one presentation.</p>
 Additional fee	<p>+ 17 000 EURO</p> <p>For each additional strength or pharmaceutical form including one presentation submitted at the same time as the initial application for authorisation.</p>
 Additional fee	<p>+ 8 600 EURO</p> <p>For each additional presentation of the same strength and pharmaceutical form, submitted at the same time as the initial application for authorisation.</p>

 Basic fee	<p>146 200 EURO</p> <p>For applications for a marketing authorisation for similar biological products authorised pursuant to Article 19 of Regulation (EU) 2019/6. This fee is for a single strength associated with one pharmaceutical form and one presentation.</p>
 Additional fee	<p>+ 17 000 EURO</p> <p>For each additional strength or pharmaceutical form including one presentation submitted at the same time as the initial application for authorisation.</p>
 Additional fee	<p>+ 8 600 EURO</p> <p>For each additional presentation of the same strength and pharmaceutical form, submitted at the same time as the initial application for authorisation.</p>

The number of target species applied for does not impact on the fee.

2.1.2.4. Reduced fee – Immunologicals

	43 300 EURO	For a single strength associated with one pharmaceutical form and one presentation.
Basic fee		
	+ 8 600 EURO	For each additional strength or pharmaceutical form including one presentation submitted at the same time as the initial application for authorisation.
Additional fee		
	+ 8 600 EURO	For each additional presentation of the same strength and pharmaceutical form, submitted at the same time as the initial application for authorisation.
Additional fee		

The number of target species applied for does not impact on the fee.




Note: Refer to section [A.2.1.2.](#) in the annex for examples of the determination of fees for applications for marketing authorisation.

2.1.3. Deleted**2.1.4. Variation to a marketing authorisation requiring assessment (VRA)**



When a change affecting different strengths, pharmaceutical forms and/or presentations is introduced through a variation (e.g. to add of a new manufacturing site for active substance, to update the safety data on the product information), the corresponding fee for the variation covers all authorised strengths, pharmaceutical forms and presentations of a given medicinal product (= per main authorisation number) except for Level 1 VRA for which additional fees are foreseen for additional presentation or strength/potency included in the same variation.

The following fees apply to variations requiring assessment that are notified or applied for individually in an application⁵.





2.1.4.1. Variation requiring assessment introducing changes of active substance(s), strength, pharmaceutical form, route of administration or food-producing target species (Level 1 VRA)

	43 300 EURO	For a variation requiring assessment that introduces changes of route of administration or food-producing target species
Basic fee		
	39 100 EURO	For a variation requiring assessment that introduces changes of active substance(s), strength or pharmaceutical form, i.e. for which no clinical or non-clinical data are submitted or no cross-references to previously submitted clinical or non-clinical data are made by the marketing authorisation holder
Basic fee (Level II)		
	10 700 EURO	For immunological veterinary medicinal products for a variation requiring assessment that introduces changes of active substance(s), strength or
Basic fee		



⁵ Published [Guidance on the details of the classification of variations requiring assessment according to Article 62 of Regulation \(EU\) 2019/6 for veterinary medicinal products and on the documentation to be submitted pursuant to those variations](#)

(Level III)	pharmaceutical form, i.e. for which no clinical or non-clinical data are submitted or no cross-references to previously submitted clinical or non-clinical data are made by the marketing authorisation holder
	+ 10 700 EURO
Additional fee	For each additional strength/potency of the same pharmaceutical form of the same variation application including one presentation submitted at the time of the variation application. This fee shall also cover one or more target species associated with that pharmaceutical form.
	+ 8 600 EURO
Additional fee	For each additional presentation of the same strength/potency of the same pharmaceutical form of the same variation application submitted at the same time of the variation application.

2.1.4.2. Variation requiring assessment introducing safety, efficacy or pharmacovigilance changes (Level 2 VRA)

	51 800 EURO
Basic fee (Level I)	For variations requiring assessment that introduce changes to safety, efficacy or pharmacovigilance, which follow an extended (90 days) or a standard (60 days) timetable (except for scope G.I.18 as set out in basic fee level IV below)
	8 600 EURO
Basic fee (Level II)	For immunological veterinary medicinal products for variations requiring assessment that introduce changes to safety, efficacy or pharmacovigilance, which follow an extended (90 days) or a standard (60 days) timetable
	13 000 EURO
Basic fee (Level III)	For each of the third and subsequent variation introducing changes specified for the Level 2 VRA basic fee level I above that is grouped in a single application made under the terms of Article 64 of Regulation (EU) 2019/6. For each of the third and subsequent variation introducing changes specified for the Level 2 VRA basic fee level I above in the case of a worksharing application under Article 65 of Regulation (EU) 2019/6.
	8 600 EURO
Basic fee (Level IV)	For variations requirement assessment with scope G.I.18 referred to in the guidance on the details of the classification of variations requiring assessment according to Article 62 of Regulation (EU) 2019/6 for veterinary medicinal products and on the documentation to be submitted pursuant to those variations

2.1.4.3. Variation requiring assessment introducing quality changes (Level 3 VRA)

	39 100 EURO
Basic fee (Level I)	For variations requiring assessment introducing quality changes only, which follow a standard (60 days) timetable, i.e. all amendments to the chemical, pharmaceutical and biological documentation, for which no clinical or non-clinical data are submitted or no cross-references to previously submitted clinical or non-clinical data are made by the marketing authorisation holder
	8 600 EURO
	For immunological veterinary medicinal products for variations requiring

Basic fee (Level II) assessment introducing quality changes only, which follow a standard (60 days) timetable, i.e. all amendments to the chemical, pharmaceutical and biological documentation, for which no clinical or non-clinical data are submitted or no cross-references to previously submitted clinical or non-clinical data are made by the marketing authorisation holder



13 000 EURO

Basic fee (Level III)

For each of the third and subsequent variation introducing changes specified for the Level 3 VRA basic fee level I above that is grouped in a single application made under the terms of Article 64 of Regulation (EU) 2019/6.

For each of the third and subsequent variation introducing changes specified for the Level 3 VRA basic fee level I above in the case of a worksharing application under Article 65 of Regulation (EU) 2019/6.

2.1.4.4. Variation requiring assessment (VRA) reduced timetable (Level 4 VRA)



8 600 EURO

Basic fee (Level I)

For all variations requiring assessment which follow a reduced (30 days) timetable

2.1.5. Grouping and worksharing procedures for variations

2.1.5.1. Grouping of variations requiring assessment submitted under the terms of Article 64 of Regulation (EU) 2019/6

- The applicable fees as specified in sub-section 2.1.4 shall be payable for each individual variation to a marketing authorisation that is grouped in a single application made under the terms of Article 64 of Regulation (EU) 2019/6.
- The applicable level I and level II basic fees specified in sub-section 2.1.4.2 and 2.1.4.3 above are payable for the first and second VRA respectively when both levels of fees are applicable to VRA in the same grouping.
- Subsequent variations in a grouping shall be similarly charged the applicable fees as specified above.
- The same marketing authorisation holder also means several marketing authorisation holders that are linked through a parent company. The fee for the grouping shall be payable by the marketing authorisation holder applying for the grouping procedure.
- Where any variation requiring assessment included in a grouping are found not to be valid and the remainder are validated positively, the applicable fees as specified above shall be payable for each of the positively validated variations requiring assessment.

2.1.5.2. Worksharing procedure under the terms of Article 65 of Regulation (EU) 2019/6

- The applicable fee specified in section 2.1.4 is payable for each individual variation to one of the centralised marketing authorisations owned by the same holder, where more than one centralised marketing authorisation is included in the worksharing application, or to the single centralised marketing authorisation included in the worksharing application, as applicable.

- The applicable level I and level II basic fees specified in 2.1.4.2 and 2.1.4.3 above are payable for the first and second VRA respectively, when both levels of fees are applicable to variations in the same worksharing procedure.
- The administrative fee shown in the table below is additionally payable for each individual VRA to each of the other centralised marketing authorisation(s) owned by the same holder included in the same worksharing application, if applicable.
- No fee is payable to the Agency for any national marketing authorisations included in the same worksharing application.
- The same marketing authorisation holder also means several marketing authorisation holders that are linked through a parent company. The fee for the worksharing procedure shall be payable by the marketing authorisation holder applying for the procedure.

Variation type	Worksharing fees for one centralised marketing authorisation	Worksharing administrative fees for second and subsequent marketing authorisations	
		Excluding applications on usage patent grounds	Applications on usage patent grounds ⁶
VRA Level 2 basic fee level I	51 800 EURO	4 290 EURO	2 480 EURO
VRA Level 3 basic fee level I	39 100 EURO		
VRA Level 2 and 3 basic fee level III	13 000 EURO		
VRA level 2 and 3 basic fee level II	8 600 EURO		
VRA Level 4	8 600 EURO	1 450 EURO	

Where any variations included in a worksharing procedure are found not to be valid and the remainder are validated positively, the applicable fees as specified above shall be payable for each of the positively validated variations.

2.1.6. Deleted

2.1.7. Inspection



26 200 EURO

Basic fee (Level I)

For each inspection inside or outside the European Union; for inspections outside the European Union, travel expenses shall be charged extra on the basis of actual cost.



13 000 EURO

Basic fee (Level II)

For each inspection inside or outside the European Union cancelled due to the withdrawal of the application; or changes to the manufacturing arrangements made by the manufacturer or changes made by the applicant/MAH that necessitate a cancellation of the inspection before the inspection is carried out.

⁶ These fees are applicable for as long as the concerned marketing authorisation is affected by usage patent(s) pertaining to indication(s) and/or dosage form(s). Refer to section 4.3 for further details on fee exemptions.

Note: Refer to section [A.1.1.7.1.](#) in the annex for examples of the determination of fees for good manufacturing practice (GMP) inspections and to section [A.1.1.7.2.](#) in the annex for examples of the determination of fees for good clinical practise (GCP).

2.1.8. Transfer of a marketing authorisation



8 600 EURO

Basic fee

For all authorised presentations of the medicinal product.

2.1.9. Maintenance of a marketing authorisation – Annual fee



41 500 EURO

Basic fee
(Level I)

For each marketing authorisation of a veterinary medicinal product. This fee covers all authorised presentations of the medicinal product.



20 600 EURO

Basic fee
(Level II)

For each marketing authorisation of a similar biological veterinary medicinal products authorised pursuant to Article 19 of Regulation (EU) 2019/6. This fee covers all authorised presentations of the medicinal product.



10 200 EURO

Basic fee
(Level III)

For each marketing authorisation of a veterinary medicinal product authorised pursuant to Articles 18, 19 (except for the medicinal products covered in basic fee level II above), 21 and 23 of Regulation (EU) 2019/6. This fee covers all authorised presentations of the medicinal product.

2.1.10. Referral



51 800 EURO

Basic fee

For the procedures laid down in Article 82 of Regulation (EU) 2019/6 initiated by the applicant of a marketing authorisation or the marketing authorisation holder.

Note: Where more than one applicant of marketing authorisations or holder of existing marketing authorisations are concerned by the procedures referred to in the first subparagraph, the applicants or holders may be grouped for the purpose of the payment of one single referral fee. If, however, the same procedure concerns more than 10 different applicants or holders, the fee will be charged by the application of the above-mentioned referral fee.

2.2. Maximum residue limits (MRLs)



86 000 EURO

Basic fee

For an application to set an initial MRL for a given substance intended to be used in veterinary medicinal products or in biocidal products used in animal husbandry.



26 200 EURO

Basic fee

For each application to modify or extend an existing MRL, as included in Table 1 of the Annex to Regulation (EEC) No 470/2009.

MRL fees shall be deducted from the fee payable for an application for marketing authorisation or an application to add a food-producing target species to a marketing authorisation for the medicinal product containing the substance for which an MRL has been set where such applications are submitted by the same applicant. However, this deduction may in total be no more than one half of the fee to which it applies.

2.3. Scientific advice

2.3.1. Definitions

The following definitions shall apply for the determination of fees for scientific advice requests.

Quality development: Chemical, pharmaceutical and biological testing.

Safety development: Toxicological and pharmacological tests.

Clinical development: Studies in animal patients, including clinical pharmacological trials designed to determine the efficacy and safety of the product.

Initial request: First request for scientific advice or protocol assistance introduced in relation to the submission of an application for marketing authorisation or a variation, whatever the authorisation phase (pre- or post-authorisation).

Follow-up to initial request: Any subsequent request falling within the same therapeutic indication and initial area(s) as the initial request, (area meaning quality, preclinical and/or clinical development including pharmacovigilance/risk management aspects).

2.3.2. Initial request for scientific advice



17 000 EURO

Basic fee
(Level I)

For **initial** requests on:

- quality development, or
- safety development, or
- bioequivalence studies for generic medicinal products, or
- new MRL, or
- preliminary risk profile



26 200 EURO

Basic fee
(Level II)

For **initial** requests on:

- clinical development, or
- quality and safety development, or
- quality and bioequivalence studies for generic medicinal products.



51 800 EURO

Basic fee
(Level III)

This fee corresponds to an initial request for scientific advice (SA) on:

- quality and safety and clinical development, or
- quality and clinical development, or
- safety and clinical development.

2.3.3. Follow-up request for scientific advice



13 000 EURO

Basic fee
(Level I)

For **follow-up** to the initial request on:

- quality development, or
- safety development
- bioequivalence studies for generic medicinal products, or
- new MRL, or
- preliminary risk profile



17 000 EURO

Basic fee
(Level II)

For **follow-up** to the initial request on:

- clinical development, or
- quality and safety development, or
- quality and bioequivalence studies for generic medicinal products.



26 200 EURO

Basic fee
(Level III)

For **follow-up** to the initial request on:

- quality and safety and clinical development, or
- quality and clinical development, or
- safety and clinical development.

2.4. Scientific services

A scientific service fee shall apply where an application is made for any scientific advice or opinion by a scientific committee, which is not covered by Articles 3 to 7 or by Article 8(1) of Council Regulation (EC) No 297/95.

2.4.1. Assessment on requirement for full MRL evaluation



26 200 EURO

Basic fee

Assessment on whether a full MRL evaluation is required or not for a chemical-unlike biological substance.

2.4.2. Classification for limited market and eligibility for Article 23 of Regulation (EU) 2019/6



13 000 EURO

Basic fee

For a request for classification of a veterinary medicinal product as intended for a limited market according to Article 4(29) and as eligible for authorisation according to Article 23 of Regulation (EU) 2019/6.

2.4.3. Certification of compliance with European Union legislation for vaccine antigen master files (VAMF)



8 600 EURO

Basic fee

For the review of the VAMF and its certification when it is submitted in parallel and within the submission of a new application containing the named antigen within the centralised procedure.



+ 8 600 EURO

Additional fee

For each VAMF application submitted simultaneously submitted simultaneously in the context of the same new marketing authorisation application (e.g. multivalent vaccines) up to a maximum of EUR 26 200.



17 000 EURO

Basic fee

For the review of the VAMF and its certification where the initial data have been previously evaluated within the centralised, decentralised or mutual recognition procedure.



8 600 EURO

Basic fee

For review and certification of a variation requiring assessment to the VAMF in accordance with Commission Regulation (EU) 2019/6 and section 2.1.5 of this explanatory note.

2.4.4. Certification of compliance with European Union legislation for vaccine platform technology master files (vPTMF)



8 600 EURO

Basic fee

For the review of the vPTMF and its certification when it is submitted in parallel and within the submission of a new application containing the named platform within the centralised procedure.



17 000 EURO

Basic fee

For the review of the vPTMF and its certification for a platform in vaccine(s) already authorised via the centralised, decentralised or mutual recognition procedure



8 600 EURO

Basic fee

For review and certification of a variation requiring assessment to the vPTMF in accordance with Commission Regulation (EU) 2019/6 and section 2.1.5 of this explanatory note.



Administrative fees

3. Administrative fees

3.1. Negative validation



3 750 EURO

Basic fee

For an application that has been found not to be valid, an administrative fee for the validation of the application shall be charged.

This fee is for all negative validations except for grouping or worksharing applications.



3 750 EURO

Basic fee

For a grouping or worksharing application where all extensions/variations in the application have been found not to be valid.

If individual extensions/variations in an application are found not to be valid but the remainder are validated positively, no administrative fee shall be charged for the invalid extensions/variations.

3.2. Certificate of a medicinal product

3.2.1. Definitions

A set of certificates: a maximum of six identical original certificates for a medicinal product with a distinct marketing authorisation number, addressed to the same importing country, issued in the same official language of the European Union and having identical annexes.

Standard procedure: procedure for issuing certificates of medicinal product within 10 working days.

Urgent procedure: procedure for issuing certificates of medicinal product within 2 working days.

3.2.2. Standard procedure



360 EURO

Basic fee
(Level I)

For each request for certificates, including one set of certificates.



+ 180 EURO

Additional fee

For each additional set of certificates included in the same request.

3.2.3. Urgent procedure



1 080 EURO

Basic fee
(Level II)

For each request for certificates, including one set of certificates.



+ 540 EURO

Additional fee

For each additional set of certificates included in the same request.

3.2.4. Withdrawal of request for certificates



360 EURO

Basic fee

When a request for certificates is withdrawn by the requester following confirmation by the Agency of the start of the procedure.

Note: Refer to section [A.3.2.](#) in the annex for examples of the determination of fees for certificates of medicinal product.

3.3. Notification of parallel distribution



3 750 EURO

Initial notification of parallel distribution

For each EU presentation of a medicinal product for one Member State of destination having one or more official languages or for several Member States of destination having the same official language.

This fee covers any subsequent safety update notification relating to the initial notification triggered by a safety update adopted by the Committee for Medicinal Products for Human Use or the Committee for Veterinary Medicinal Products, which is identified and communicated by the Agency to the parallel distributor.



720 EURO

Annual update notification, **manual** check

For all changes made in one year from the anniversary of the initial notification to all the EU presentations belonging to the same pharmaceutical form of the same medicinal product for one Member State of destination having one or more official languages, or for several Member States of destination having the same official language.

This fee applies when a report generated by text comparison software has not been provided or has not met the specific conditions laid down by the Agency for the acceptance of the report.



350 EURO

Annual update notification, **automated** check

For all changes made in one year from the anniversary of the initial notification to all the EU presentations belonging to the same pharmaceutical form of the same medicinal product for one Member State of destination having one or more official languages, or for several Member States of destination having the same official language.

This fee applies when a report generated by text comparison software has been provided and has met the specific conditions laid down by the Agency for the acceptance of the report.



720 EURO

Notification of changes, **manual** check

For each notification of changes that is not submitted as part of the annual update notification and is not a safety update.

This fee applies for each EU presentation of a medicinal product for one Member State of destination having one or more official languages, or for several Member States of destination having the same official language.

This fee applies when a report generated by text comparison software has not been provided or has not met the specific conditions laid down by the Agency for the acceptance of the report.



350 EURO

Notification of changes, **automated** check

For each notification of changes that is not submitted as part of the annual update notification and is not a safety update.

This fee applies for each EU presentation of a medicinal product for one Member State of destination having one or more official languages, or for several Member States of destination having the same official language.

This fee applies when a report generated by text comparison software has been provided and has met the specific conditions laid down by the Agency for the acceptance of the report.



3 750 EURO

Notification of bulk changes

For one or more changes that affect all of a parallel distributor's initial notifications, at any point in time after the approval of the initial notification. The scope(s) of the changes are limited to: a change in the details of a parallel distributor ('Change of name and/or address' or 'Reassignment of notices for Parallel Distribution'), addition or deletion of a re-packager, and/or a change in the name and/or address of a re-packager.

3.4. Worksharing procedures for variations to marketing authorisations

Refer to section 1.1.5 for details of administrative charges applicable to worksharing procedures under the terms of Article 20 of Commission Regulation (EC) No 1234/2008 and 2.1.5 for details of administrative charges applicable to worksharing procedures under the terms of Article 65 of Regulation (EU) 2019/6.



Fee determination and payment

4. Fee determination and payment

4.1. Legal requirements

Article 10 of Council Regulation (EC) No 297/95 stipulates the dates when fees are due and when they are payable.

Fee due date

The fee due date is the date when the Agency determines the total fee amount that is due for an application or service. The applicant's or marketing authorisation holder's obligation to pay that fee starts on that date.

The fee due date corresponds to:

- The date of administrative validation of an application; or
- The date of the start of a procedure (when there is no administrative validation), e.g. for type IA variation fees, transfer fees, Article 30(1) or 31 referrals; or
- The date of the anniversary of the notification of the marketing authorisation decision, for annual fees; or
- The date of the receipt of the inspection report, for inspection fees.

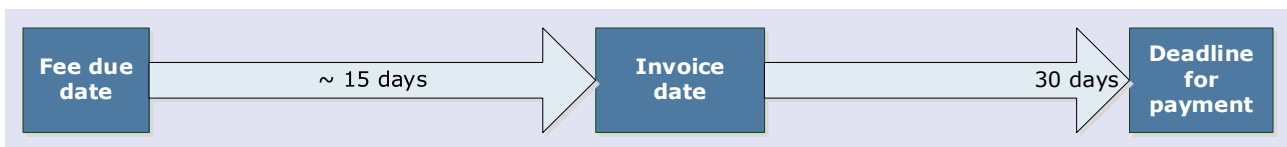
Note: For a change of applicant during an ongoing initial marketing authorisation application, any fee invoiced since validation (i.e. fee for initial marketing authorisation and pre-authorisation inspection fee) will be credited to the original applicant and re-invoiced to the new applicant. This includes changes, if any, relating to micro, small or medium-sized enterprise applicants and orphan medicinal product designation.

Payment of fees

The Agency's terms and conditions for payment of fees stipulate that an invoice is payable 30 days from the invoice date. That represents the deadline for the applicant to settle the payment.

In summary, the procedure shown in Figure 1 applies.

Figure 1. Procedure for fee determination and payment



Deferred fees

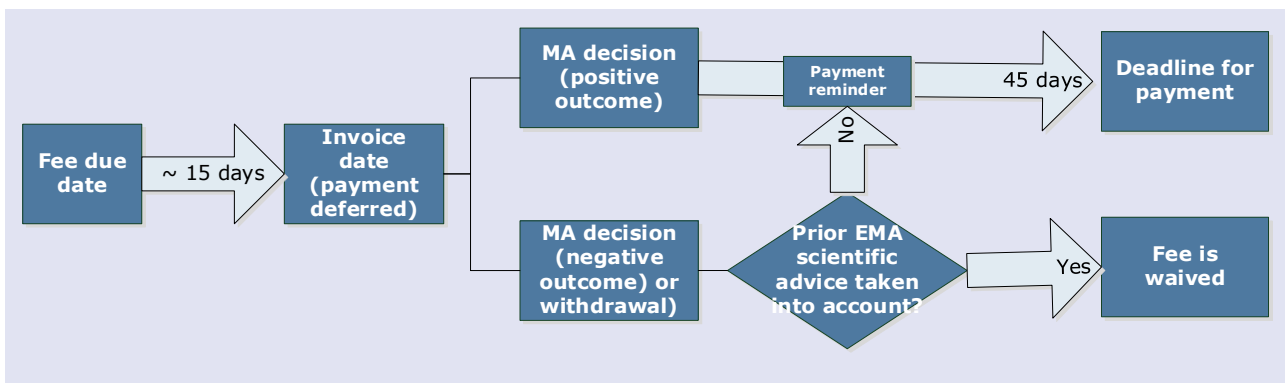
Small or medium-sized enterprise (SME)

When the applicant is an SME, payment of the fee for a marketing authorisation (MA) application and the fee for inspections undertaken as part of the assessment of a MA application is deferred until the final decision (positive or negative outcome) on the MA is issued or the application is withdrawn. The invoice issued at the time the fee is due informs the applicant that the fee is deferred. A reminder is issued by the Agency as soon as there is a decision or withdrawal; the fee is payable within 45 days of the date of the notification of the final decision on the MA or of the withdrawal of the application.

A conditional fee exemption of the fee for the examination of a MA application is given where the scientific advice provided by EMA was taken into account by the applicant for the development of the product and a MA is not granted (due to negative outcome or withdrawal of the MA application). To benefit from a conditional fee exemption, the required criteria shall be valid on the fee due date. A subsequent change to the SME status (i.e. expiration of SME status) after the fee due date will not be taken into account.

The procedures are summarised in Figure 2.

Figure 2. Procedure for fee determination and payment by SME



Core dossier submission

The payment of the fee for an application for a MA of a medicinal product to be used in a human pandemic situation, so-called core dossier submission, is deferred until the pandemic situation is duly recognised, either by the World Health Organisation or the European Union, or up to 5 years from the due date, whichever comes first. Nevertheless, the deferral can come to an end earlier on withdrawal prior to an opinion on the MA application, or if a negative Commission Decision is issued. A reminder is issued by the Agency as soon as the deferral comes to an end and the fee is payable immediately.

4.2. Criteria for fee determination

The criteria that are taken into account by the Agency when determining the fee that is due are the ones that apply by the fee due date. These include:

- The applicant's status (e.g. valid SME status assigned by the Agency);
- The product type (e.g. advanced therapy medicinal product, designated orphan product);
- The product status (e.g. in the first year following grant of a MA); and
- The procedure type.

Any change to the applicable criteria after the fee due date is not taken into account and will not influence the total fee due that has been determined.

4.3. Established fee exemptions

An applicant qualifies for fee exemptions and incentives specified in Chapter 5 provided that the required criteria apply on the fee due date. Any criterion that is no longer valid on the fee due date, e.g. expired SME status, cannot be taken into consideration by the Agency.

Applicants should ensure that they meet all the required criteria at the time of submission of an application or a request for a service.

4.4. Request for ad hoc fee reduction

Applicants may request the Executive Director to grant an ad hoc fee reduction under the provisions of the first paragraph of Article 9 of Council Regulation (EC) No 297/95.

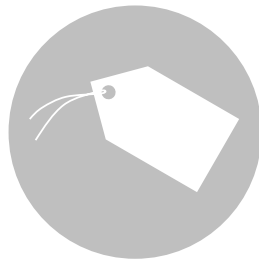
The applicant is required to provide sufficient justification to prove that the request is made (1) in exceptional circumstances and (2) for imperative reasons of public or animal health. The request will be considered by the Executive Director, who will consult the competent scientific committee and will decide each request on a case-by-case basis.

In view of the administrative procedure that has to be followed and the mandatory scientific consultation, applicants are required to make their request in a letter to the Executive Director at least two months before the date of submission of the relevant application or two months before the anniversary of the European Birth date of the Marketing Authorisation for Annual fees. The applicant should cite Article 9, paragraph 1 of Council Regulation (EC) No 297/95 and provide details of the product, procedure type and applicable fee, and the reason(s) for the request that justify exceptional circumstances and imperative reasons of public or animal health.

Applicants are advised that late requests may not be processed in time and may not be taken into consideration when determining the fee.

Applicants are also advised that requests submitted after the receipt of an invoice will not be considered.

Further information on fees and fee exemptions is available on the Agency's [website](#).



Fee exemptions

5. Fee exemptions

Where an applicant could, in respect of the same fee, benefit from more than one category of fee reduction or incentive (e.g. advanced therapy medicinal product, orphan medicinal product, micro, small or medium sized enterprises) the provisions which are the most favourable to the applicant would apply. The applicant can contact the Agency prior to submission of the application for confirmation of the applicable fee.

5.1. Micro, small or medium-sized enterprise

5.1.1. Definitions

Pursuant to Regulation (EC) No 2049/2005, Applicants that meet the definition of a micro, small or medium-sized enterprise are eligible for fee incentives from the Agency. Companies developing medicinal products for human and/or veterinary use can benefit from fee reductions, exemptions and/or deferrals, as applicable, under Articles 5 to 9 of Regulation (EC) No 2049/2005. Applicants must be established in the EEA and fulfil the definition of a micro, small or medium-sized enterprise as set out in Commission Recommendation 2003/361/EC of 6 May 2003.

5.1.2. Fee incentives

Scientific advice	90% reduction to the total applicable fee for non-orphan medicinal products
	100% reduction to the total applicable fee for designated orphan medicinal products (see section 5.2.3)
	100% reduction to the total applicable fee for products eligible to the PRIME scheme ^a
	90% reduction to the total applicable fee for veterinary medicinal products
Inspection (pre-authorisation)	90% reduction to the total applicable fee
	Deferral of total applicable fee
Application for a marketing authorisation	Deferral of total applicable fee
	Conditional fee exemption
Scientific services ^b	90% reduction to the total applicable fee for non-orphan medicinal products
	100% reduction to the total applicable fee for designated orphan medicinal products (see section 5.2.3)
Establishment of maximum residue limit for a veterinary medicinal product	90% reduction to the total applicable fee

Extension or modification of maximum residue limit for a veterinary medicinal product	90% reduction to the total applicable fee
Administrative services (excluding parallel distribution)	100% reduction to the total applicable fee
Inspection (post-authorisation)	90% reduction to the total applicable fee
Post-authorisation activities ^c	100% reduction to the total applicable fee, in the case of micro enterprises
	40% reduction to the total applicable fee, in the case of small or medium-sized enterprises

^a Fee reduction is restricted to the development in the indication for which eligibility to the PRIME scheme was accepted.

^b The fee reduction to the scientific service fee shall apply also to

- consultations on medical devices when the manufacturer has been assigned SME status by the Agency.
- veterinary medicinal product procedures with regards to Regulation (EU) 2019/6 classified as scientific service in accordance with heading 8 of Annex II to the Fee Regulation Implementing Rules.

^c Defined as: extension of a marketing authorisation (medicines for human use); type-IA, type-IB or type-II variation (medicines for human use); renewal of a marketing authorisation (medicines for human use); transfer of a marketing authorisation from a micro, small or medium-sized enterprise; annual fee; referral procedure laid down in Article 30(1) or the first sub-paragraph of Article 31(1) of Directive 2001/83/EC initiated by the marketing authorisation holder).

5.2. Orphan medicinal products

5.2.1. Definitions

Total or partial exemptions from the payment of fees for applications for designated orphan medicinal products for human use shall be granted as laid down in a decision of the Executive Director on the use of the special contribution from the European Union, provided for by Article 7(2) of Regulation (EC) No 141/2000, reflecting the advice of the Committee for Orphan Medicinal Products.⁷

Applicants from the academic sector must be established in the EEA and fulfil the definition of public or private higher education establishments awarding academic degrees, public or private non-profit research organisations whose primary mission is to pursue research, or international European interest organisations as set out in Commission Regulation (EU) No 1290/2013 of 11 December 2013.

Applicants should not be financed or managed by private profit organisations in the pharmaceutical sector ("PPO"), nor should have they concluded any operating agreements with any PPO concerning their sponsorship or participation to the specific research project for which a fee exemption is sought.

These requirements shall be evidenced by a supporting declaration and documents which will be checked by the EMA upon receipt of a scientific advice request.

⁷ Executive Decision of 19 June 2020 (EMA/135645/2020)

5.2.2. Fee incentives for applicants other than micro, small and medium sized enterprises

Protocol assistance (non-paediatric-related*) other than for academia	75% reduction to the total applicable fee
Protocol assistance for academia	100% reduction to the total applicable fee
Protocol assistance (paediatric-related*)	100% reduction to the total applicable fee
Inspection (pre-authorisation)	100% reduction to the total applicable fee
Application for a marketing authorisation	10% reduction to the total applicable fee

* Paediatric-related protocol assistance is restricted to the development of an orphan medicinal product for the paediatric population, where the advice requested does not include the adult population.

5.2.3. Fee incentives for micro, small and medium sized enterprises

Protocol assistance	100% reduction to the total applicable fee
Scientific services**	100% reduction to the total applicable fee (see section 5.1.2)
Inspection (pre-authorisation)	100% reduction to the total applicable fee
Application for a marketing authorisation	100% reduction to the total applicable fee
Post-authorisation activities, including annual fees, during the first year after marketing authorisation	100% reduction to the total applicable fee
Inspection (post-authorisation)**	90% reduction to the total applicable fee (see section 5.1.2)

** Fee reductions for scientific services and post-authorisation inspections are not funded by the special contribution from the European Union for designated orphan medicinal products but are provided for by Article 7 of Regulation (EC) No 2049/2005 on SMEs.

5.3. Multiple applications on usage patent grounds

5.3.1. Definitions

The full or partial exemptions from payment of fees described below are applicable for as long as the concerned marketing authorisation is affected **by usage patent(s)** pertaining to indication(s) and/or dosage form(s).

5.3.2. Fee incentives for an application for a marketing authorisation

5.3.2.1. Reduced fee (human medicinal products)

Second and each subsequent multiple application submitted under Articles 10(1) and 10(3) of Directive 2001/83/EC	25 700 EURO
Second and each subsequent multiple application submitted under Article 10(4) of Directive 2001/83/EC	42 900 EURO
Additional strengths, pharmaceutical forms and presentations submitted at the same time as the aforementioned applications	100% reduction to the total applicable fee

5.3.2.2. Reduced fee (veterinary medicinal products)

Second and each subsequent multiple application submitted under Articles 18 and 19 (except products specified below) of Regulation (EU) 2019/6 solely on usage patent grounds	17 000 EURO
Second and each subsequent multiple application submitted for similar biological veterinary medicinal products submitted under Articles 19 of Regulation (EU) 2019/6 solely on usage patent grounds	31 000 EURO
Second and each subsequent multiple application submitted for veterinary immunological medicinal products solely on usage patent grounds	17 000 EURO
Additional strengths, pharmaceutical forms and presentations submitted at the same time as the aforementioned applications	100% reduction to the total applicable fee

5.3.3. Fee incentives for post-authorisation activities for a second and for each subsequent multiple application

5.3.3.1. Reduced fee (human medicinal products)

Extension of a marketing authorisation	24 700 EURO
Additional strengths and presentations of the same pharmaceutical form submitted at the same time as the aforementioned application	100% reduction to the total applicable fee

Type-IA variation to a marketing authorisation. This fee shall only apply in the case of grouping of the same type-IA variations to the terms of multiple marketing authorisations on usage patent grounds owned by the same holder (as defined in Article 7(2)(a) of Commission Regulation (EC) No 1234/2008). The applicable fee shall be payable for each individual type-IA variation relating to the second and each of the subsequent multiple marketing authorisations in the grouping	720 EURO
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Worksharing procedure under the terms of Article 20 of Commission Regulation (EC) No 1234/2008. Administrative fees for variations to multiple centralised marketing authorisations on usage patent grounds	Refer to section 1.1.5.2.
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Renewal of a marketing authorisation	3 310 EURO
Additional strengths associated with a pharmaceutical form submitted at the same time as the aforementioned application	100% reduction to the total applicable fee

Annual fee for a marketing authorisation granted under Articles 10(1) and 10(3) of Directive 2001/83/EC	6 000 EURO
Annual fee for a marketing authorisation granted under Article 10(4) of Directive 2001/83/EC	11 800 EURO

5.3.3.2. Reduced fee (veterinary medicinal products)

Second and each subsequent multiple application submitted solely on usage patent grounds for variation requiring assessment specified in sub-section 2.1.4.1 of marketing authorisations granted under Articles 18 and 19 of Regulation (EU) 2019/6	8 660 EURO
Additional strengths and presentations of the same pharmaceutical form submitted at the same time as the aforementioned application	100% reduction to the total applicable fee

Worksharing procedure under the terms of Article 65 of Regulation (EU) 2019/6. Administrative fees for variations to multiple centralised marketing authorisations on usage patent grounds	Refer to section 2.1.5.2.
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Annual fee for a marketing authorisation granted under Articles 18 and 19 (except products specified below) of Regulation (EU) 2019/6	2 060 EURO
Annual fee for a marketing authorisation granted under granted under Article 19 of Regulation (EU) 2019/6 for similar biological veterinary medicinal products	4 290 EURO

5.4. Medicinal products for paediatric use

5.4.1. Definitions

A partial exemption from the payment of the fees laid down in the fee regulation is granted for paediatric use marketing authorisation applications (PUMAs) submitted under Article 30 of Regulation (EC) No 1901/2006 on medicinal products for paediatric use.

A total exemption from the payment of the fees laid down in the fee regulation is granted for scientific advice provided by the Agency to sponsors developing medicinal products for the paediatric population as laid down in Regulation (EC) No 1901/2006 on medicinal products for paediatric use.

5.4.2. Fee incentives for paediatric use marketing authorisations (PUMAs)

Application for a paediatric use marketing authorisation	
Inspection (pre-authorisation)	
During the first year after marketing authorisation for: <ul style="list-style-type: none"> extension of a marketing authorisation; type-IA, type-IB and type-II variations; annual fee; inspection (post-authorisation). 	

50% reduction to the total applicable fee

5.4.3. Fee incentives for scientific advice

Scientific advice on the development of a medicinal product for the paediatric population (when the advice requested does not include the adult population)	100% reduction to the total applicable fee
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5.5. Advanced therapy medicinal products

5.5.1. Definitions

Total or partial exemptions from the payment of fees for applications for advanced therapy medicinal products for human use are granted as laid down in Articles 16(2) of Regulation (EC) No 1394/2007.

5.5.2. Fee incentives for applicants other than micro, small and medium-sized enterprises

Scientific advice	65% reduction to the total applicable fee
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5.5.3. Fee incentives for micro, small and medium-sized enterprises

Scientific advice	90% reduction to the total applicable fee (see section 5.1.2)
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5.6. Core dossier medicinal products to be used in a human pandemic situation

5.6.1. Definitions

A total exemption from the payment of the fees laid down in the fee regulation is granted for the regulatory activities specified below within the framework of the submission of a core dossier for a pandemic influenza vaccine and the follow-up submission of a pandemic variation, as described in the '[Guideline on influenza vaccines – submission and procedural requirements](#)' (EMA/CHMP/VWP/457259/2014).

The following total exemptions apply until the type II pandemic variation, submitted once the human pandemic situation is duly recognised, has been authorised by the European Union.

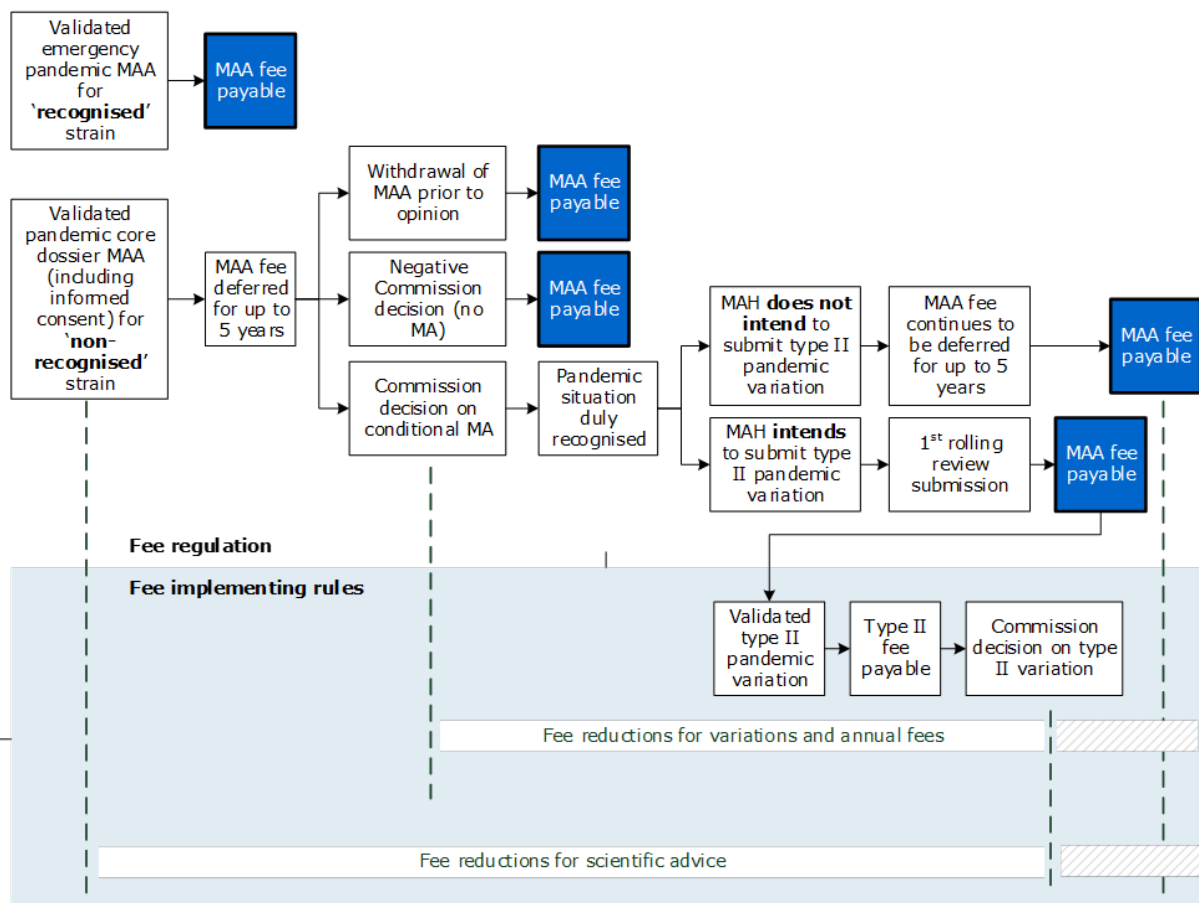
5.6.2. Fee incentives related to pandemic core dossier (including informed consent) for 'non-recognised' strain

Scientific advice	100% reduction to the total applicable fee
Post-authorisation activities including type-IA, type-IB, type-II variations (but excluding the type-II pandemic variation) and annual fee	100% reduction to the total applicable fee
Negative validation of a type-IB or type-II variation (but excluding the type-II pandemic variation)	100% reduction to the total applicable fee

- The fee for a validated pandemic core dossier marketing authorisation application (MAA), including informed consent application, for a pandemic strain that has not yet been "duly recognised, either by the World Health Organisation or by the Community" is deferred.
- If the core dossier MAA is withdrawn prior to opinion or receives a negative Commission Decision, the fee becomes payable and the deferral comes to an end.

- If the core dossier receives a positive Commission Decision, the fee shall continue to be deferred. The MAH also receives fee reductions on variations and annual fees.
- When the pandemic situation becomes duly recognised, the marketing authorisation holder (MAH) may choose **not** to submit a Type II pandemic variation. In this case the MAA fee continues to be deferred for up to 5 years from the MAA validation date. The MAH continues to receive fee reductions on variations and annual fees.
- If the MAH chooses to submit a Type II pandemic variation, the initial MAA fee becomes payable on the first submission made as part of the rolling review. When the Type II pandemic variation is validated the applicable Type II fee becomes payable. The MAH continues to receive fee reductions on variations and annual fees until the date of the Type II Commission Decision.
- Fee reductions for scientific advice are applicable to requests in the context of a pandemic core dossier and at any time from the MAA validation date up to the date of the Type II Commission Decision.
- The fee for an emergency pandemic MAA submitted when the pandemic situation has been duly recognised becomes payable on validation.

Figure 3. Schematic overview of pandemic vaccine fee reductions, deferrals and payments



5.7. Veterinary medicinal products for limited markets

5.7.1. Definitions

A total or partial exemption of the fee shall apply for veterinary medicinal products classified as eligible for consideration under Article 23 of Regulation (EU) 2019/6 as referred to in section 2.4.

5.7.2. Fee incentives

Scientific advice	50% reduction to the total applicable fee
Extension of maximum residues limit involving no assessment of data for veterinary medicinal products classified as eligible for consideration under Article 23	100% reduction to the total applicable fee
Establishment or extension of maximum residues limit requiring an assessment of data for veterinary medicinal products classified as eligible for consideration under Article 23	50% reduction to the total applicable fee

5.8. Deleted

5.9. Veterinary vaccines against certain epizootic diseases

5.9.1. Definitions

A total exemption from the payment of the fees laid down in the fee regulation is granted for certain post-authorisation activities in relation to vaccines against:

- bluetongue
- pandemic avian influenza
- foot and mouth disease
- classical swine fever

for which

- the vaccine is authorised under normal circumstances and
- the product has not been marketed within the EU/EEA at any time during the totality of the period covered by the fee.

5.9.2. Fee incentives

Annual fee	100% reduction to the total applicable fee
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5.10. Scientific advice for PRIME scheme

5.10.1. Definitions

A total exemption from the payment of the fees laid down in the fee regulation is granted for requests for scientific advice and follow-up requests submitted on products eligible to PRIME scheme for SMEs and applicants of the academic sector.

Applicants from the academic sector must be established in the EEA and fulfil the definition of public or private higher education establishments awarding academic degrees, public or private non-profit research organisations whose primary mission is to pursue research, or international European interest organisations as set out in Commission Regulation (EU) No 1290/2013 of 11 December 2013.

Applicants should not be financed or managed by private profit organisations in the pharmaceutical sector ("PPO"), nor should have they concluded any operating agreements with any PPO concerning their sponsorship or participation to the specific research project for which a fee exemption is sought.

These requirements shall be evidenced by a supporting declaration and documents which will be checked by the EMA upon receipt of a scientific advice request.

5.10.2. Fee incentives

Scientific advice	100% reduction to the total applicable fee*, in the case of micro, small or medium sized enterprises
	100% reduction to the total applicable fee*, in the case of applicants from the academic sector

* Fee reduction is restricted to the development in the indication for which eligibility to the PRIME scheme was accepted.

5.11. Medicinal products authorised under exceptional circumstances for preparedness against biological agents that might be used as weapons of bioterrorism

5.11.1. Definition

A partial exemption from the payment of the fees laid down in the fee regulation is granted for certain post-authorising activities in relation to medicinal products authorised under exceptional circumstances only for preparedness against biological agents that might be used as weapons of bioterrorism as detailed in the [EMA/CHMP Guidance document on use of medicinal products for the treatment and prophylaxis of biological agents that might be used as weapons of bioterrorism](#).⁸

Such exemption shall not apply

- to medicinal products that are also authorised for a therapeutic indication unrelated to preparedness against biological agents that might be used as weapons of bioterrorism;

⁸ CPMP/4048/01 16.01.2002 as last amended; i. e. anthrax, plague, tularaemia, smallpox, viral haemorrhaging fever, botulism, brucellosis, Q-fever, glanders and melioidosis and other infectious diseases

- to any type II variation to the terms of the marketing authorisation aiming at authorising a therapeutic indication unrelated to preparedness against biological agents that might be used as weapons of bioterrorism.

5.11.2. Fee incentives

Annual fee	50 % reduction to the total applicable fee
Type-IA, type-IB and type-II variations	50% reduction to the total applicable fee
Renewal of marketing authorisation	50% reduction to the total applicable fee
Extension of a marketing authorisation related to the preparedness	50% reduction to the total applicable fee
Transfer of a marketing authorisation	50% reduction to the total applicable fee

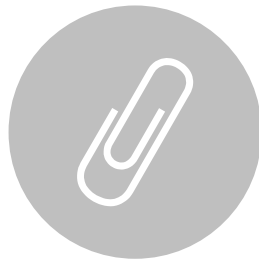
5.12. Scientific advice on clinical trial protocols pursuant to Article 16(2) of Regulation (EU) 2022/123

5.12.1. Definition

A total exemption from the payment of the fees laid down in the fee regulation is granted for certain scientific advice on clinical trial protocols for medicinal products with the potential to address a public health emergency pursuant to Article 16(2) of Regulation (EU) 2022/123.

5.12.2. Fee incentives

Scientific advice for clinical protocol	100 % reduction to the total applicable fee
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Annex

Annex

The additional information in this annex is listed using the same numbering as the corresponding sections in chapters 1 to 4. For example, section A.1.1.2. in this annex relates to section 1.1.2. in Chapter 1.

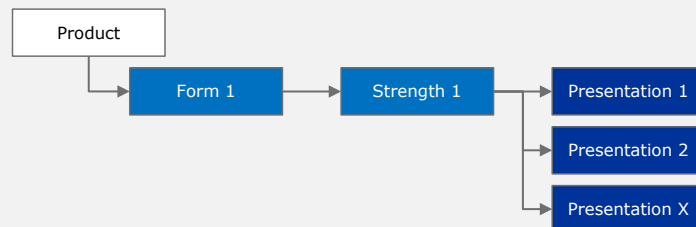
A.1. Medicinal products for human use


A.1.1.2. Examples of the determination of fees for applications for marketing authorisation


It should be noted that the calculation of the total fee for a marketing authorisation application is driven by the pharmaceutical forms, the associated strengths and the associated presentations. The examples shown below do not represent an exhaustive list.

Scenario 1:

Full dossier application for a medicinal product having one pharmaceutical form with one strength and X presentations

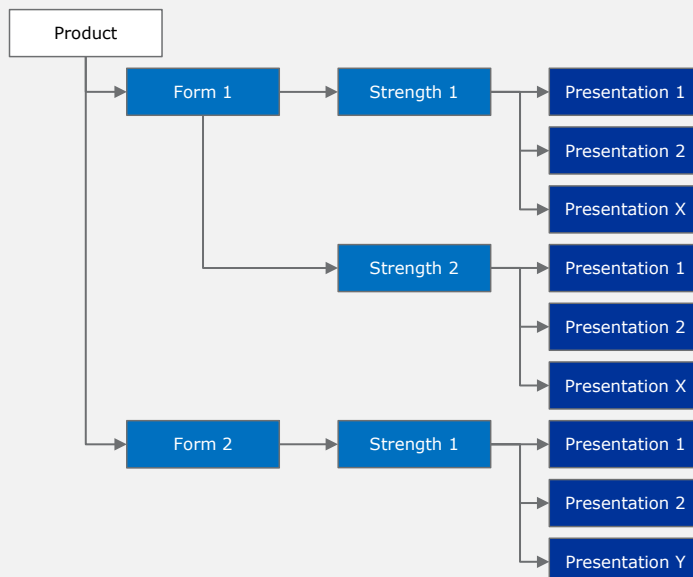



345 800 EURO
 Basic fee Includes one pharmaceutical form and one associated strength and one presentation.


+ (X-1)*8 600 EURO
 Additional fee For additional presentations associated with the single strength.

Scenario 2:

Full dossier application for a medicinal product having two pharmaceutical forms with two strengths and X presentations/ strength associated with the first form and one strength and Y presentations associated with the second form

**345 800 EURO**

Basic fee

Includes one pharmaceutical form and one associated strength and one presentation.

**+ (X-1)*8 600 EURO**

Additional fee

For additional presentations associated with the first form and strength.

**+ 34 800 EURO**

Additional fee

Second strength associated with the first form including one presentation.

**+ (X-1)*8 600 EURO**

Additional fee

For additional presentations associated with the first form and second strength.

**+ 34 800 EURO**

Additional fee

Second form including its associated strength and one presentation.

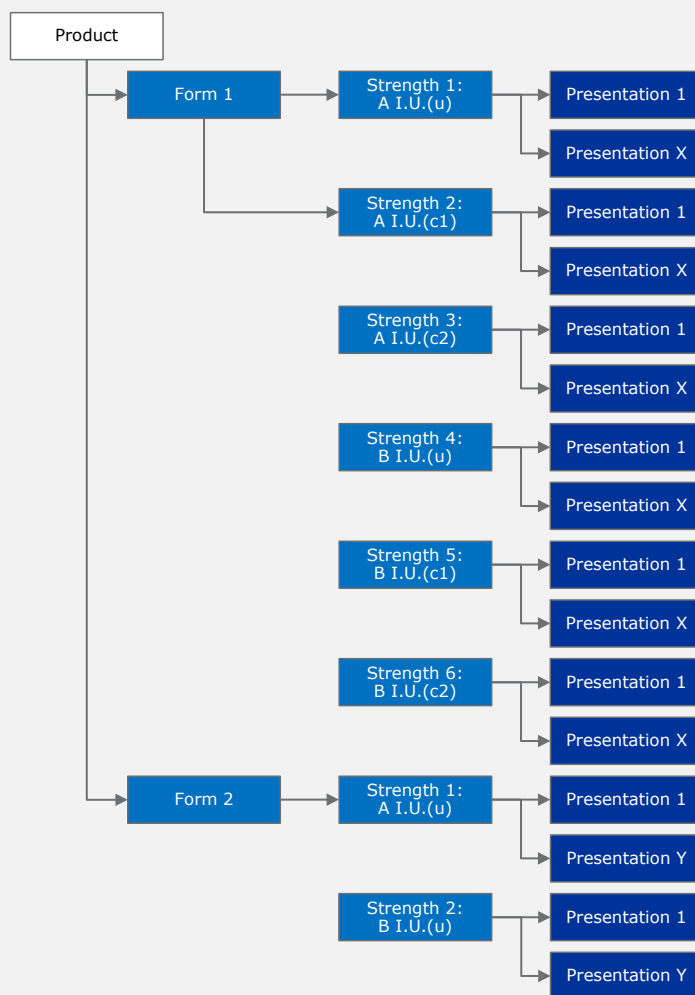
**+ (Y-1)*8 600 EURO**

Additional fee

For additional presentations associated with the second form and its strength.

Scenario 3:

Full dossier application for an insulin product having two pharmaceutical forms with six strengths (consisting of two sets of one uncombined preparation and two combination preparations (having different proportions of insulin) with insulin amounts corresponding to A I.U. and B I.U.) and X presentations/strength associated with the first form; and two strengths (of un-combined preparations with insulin amounts corresponding to A I.U. and B I.U.) and Y presentations/strength associated with the second form



(u) = un-combined insulin preparation

(c) = combination insulin preparation (refer to section 1.1.1 for definition of strength of insulin products)



345 800 EURO

Basic fee

Includes one pharmaceutical form and one associated strength and one presentation.



+ (X-1)*8 600 EURO

Additional fee

For additional presentations associated with the first form and strength.



+ 5*34 800 EURO

For second to sixth strengths associated with the first form including one

Additional fee presentation for each strength.



+ 5*(X-1)*8 600

Additional fee

Additional presentations associated with the second to sixth strengths of the first form.



+ 34 800 EURO

Additional fee

Second form including one associated strength and one presentation.



+ (Y-1)*8 600 EURO

Additional fee

For additional presentations associated with the second form and first strength.



+ 34 800 EURO

Additional fee

Second strength associated with the second form including one presentation.



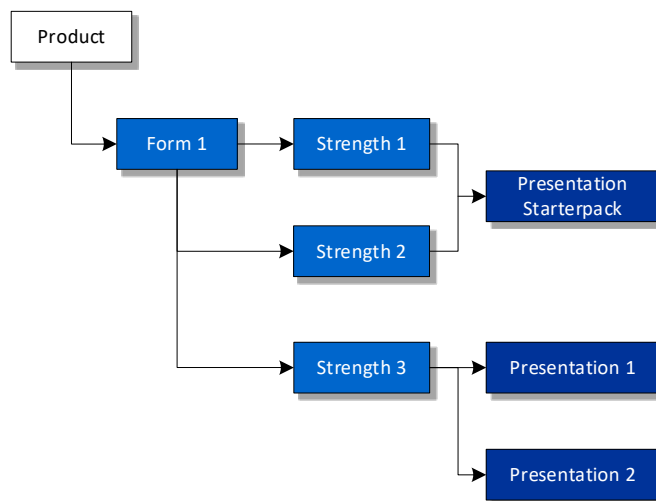
+ (X-1)*8 600 EURO

Additional fee

For additional presentations associated with the second form and second strength.

Scenario 4:

Full dossier application with 3 strengths, e.g. 100mg, 200mg and 300mg. The 100 and 200mg strengths will be packaged together in a starter pack and the 300mg strength will have two presentations. The 100 and 200mg strengths do not have additional presentations.

**345 800 EURO**

Basic fee

Includes one pharmaceutical form and one associated strength and the starter pack presentation.

**+ 2*34 800 EURO**

Additional fee

For second and third strengths associated with the first form including presentation 1 for the 3rd strength.

**+ 1* 8 600 EURO**

Additional fee

Additional presentation 2 associated with the third strength of the first form.

A.1.1.3. Examples of the determination of fees for extensions of marketing authorisation


It should be noted that the basic fee for an extension of a marketing authorisation is driven by the pharmaceutical form. The examples shown below do not represent an exhaustive list.


Scenario 1:


New pharmaceutical form with two strengths and X presentations/strength, for authorised or new route of administration (with submitted/cross-referenced clinical data)


Extension application:

- One pharmaceutical form, first strength and X presentations
- Second strength (of same new pharmaceutical form) and X presentations

 **103 800 EURO**
Basic fee For extension.

 **+ (X-1)* 8 600 EURO**
Additional fee For additional presentation fees.

 **+ 26 200 EURO**
Additional fee For additional strength fee.


 **+ (X-1)*8 600 EURO**
Additional fee For additional presentation fees.


Scenario 2:

New route of administration for authorised pharmaceutical form with two authorised strengths and X presentations/strength (with submitted/cross-referenced clinical data)

Extension application:

- Route of administration for authorised pharmaceutical form, first strength and X presentations
- Second strength (same new route of administration for same authorised pharmaceutical form) and X presentations

 **103 800 EURO**
Basic fee For extension.

 **+ (X-1)* 8 600 EURO**
Additional fee For additional presentation fees.



+ 26 200 EURO

Additional fee

For additional strength fee.



+ (X-1)*8 600 EURO

Additional fee

For additional presentation fees.

Scenario 3:

Two new strengths of same authorised pharmaceutical form and X presentations/strength (without submitted/cross-referenced clinical data)

Extension application:

- First new strength and X presentations
- Second new strength (of same authorised pharmaceutical form) and X presentations



77 900 EURO

Basic fee

For extension.



+ (X-1)*8 600 EURO

Additional fee

For additional presentation fees.



+ 26 200 EURO

Additional fee

For additional strength fee.



+ (X-1)*8 600 EURO

Additional fee

For additional presentation fees.

Scenario 4:

One new strength of each of two authorised pharmaceutical forms and X presentations/strength (without submitted/cross-referenced clinical data)

This constitutes as two extension applications

Extension application 1:

- New strength (of first authorised pharmaceutical form) and X presentations



77 900 EURO

Basic fee

For extension.





+ (X-1)*8 600 EURO

Additional fee For additional presentation fees.

Extension application 2:

- New strength (of second authorised pharmaceutical form) and X presentations

 **77 900 EURO**
For extension.
Basic fee

 **+ (X-1)*8 600 EURO**
For additional presentation fees.
Additional fee


Scenario 5:


New strength for an authorised pharmaceutical form and route of administration and X presentations AND new pharmaceutical form associated with a new strength and new route of administration and X presentations (with submitted/cross-referenced clinical data)

This constitutes as two extension applications

Extension application 1:


- New strength for an authorised pharmaceutical form and route of administration and X presentations (with submitted/cross-referenced clinical data)


 **103 800 EURO**
For extension.
Basic fee

 **+ (X-1)*8 600 EURO**
For additional presentation fees
Additional fee

Extension application 2:

- New pharmaceutical form associated with new (first) strength and new route of administration and X presentations

 **103 800 EURO**
For extension.
Basic fee

 **+ (X-1)*8 600 EURO**
For additional presentation fees
Additional fee

Scenario 6:

New pharmaceutical form associated with new route of administration for two new strengths and X presentations/strength (with submitted/cross-referenced clinical data) AND two new strengths for

an authorised pharmaceutical form and route of administration and X presentations/strength (without submitted/cross-referenced clinical data)

This constitutes as two extension applications

Extension application 1:

- New pharmaceutical form associated with new route of administration for first strength and X presentations (with clinical data)
- Second strength (same new pharmaceutical form associated with same new route of administration) and X presentations



103 800 EURO

For extension.

Basic fee



+ (X-1)*8 600 EURO

For additional presentation fees

Additional fee



+ 26 200 EURO

For additional strength fee

Additional fee



+ (X-1)*8 600 EURO

For additional presentation fees

Additional fee

Extension application 2:

- First new strength of authorised pharmaceutical form and route of administration and X presentations (without clinical data)
- Second strength (of same authorised pharmaceutical form and route of administration) and X presentations



77 900 EURO

For extension.

Basic fee



+ (X-1)*8 600 EURO

For additional presentation fees

Additional fee



+ 26 200 EURO

For additional strength fee

Additional fee



+ (X-1)*8 600 EURO

For additional presentation fees

Additional fee

•

A.1.1.4. Examples of the determination of fees for variations to a marketing authorisation introducing new presentations (i.e. EU sub-numbers)

It should be noted that the calculation of the total fee is determined by the number and type of variations (IA, IB).

Each new presentation requested (i.e. each new EU sub-number) should be submitted as a single variation attracting a separate fee (i.e. x additional presentations = x variations = x separate fees).

The type of variation (IA or IB) is determined by the currently approved range of pack sizes for each strength and pharmaceutical form. Range is defined from the smallest to the biggest approved pack size (not from '0') for the same pharmaceutical form and strength. The pack size equals to the number of units of the pharmaceutical form (e.g. tablets, sachets, ampoules, etc.) contained per outer packaging. Pack sizes not included within this range are considered to be outside of the range.

- For the addition of a new pack size where the number of units of the pack is within the range of the currently approved pack sizes for the strength and pharmaceutical form, applicants should submit a variation B.II.e.5.a).1 (IA).
- For the addition of a new pack size where the number of units of the pack is outside the range of the currently approved pack sizes for the strength and pharmaceutical form, applicants should submit a variation B.II.e.5.a).2 (IB).

In support of a timely introduction of new pack sizes to the market, EMA accepts the following approach for the introduction of various pack sizes falling outside the range within a single grouped submission. The biggest or the smallest pack size per strength outside the range should be classified as B.II.e.5.a).2 (IB). This presentation defines the new limits of the range so that any intermediate pack size for the strength and pharmaceutical form can be classified as B.II.e.5.a).1 (IA).

The example shown below does not represent an exhaustive list.

Scenario 1:

Addition of two new pack-sizes (pack-sizes 6 and 10) to each of the two strengths for a marketing authorisation with a currently approved pack-size of 4 authorised for each strength.



3 900 EURO

Basic fee

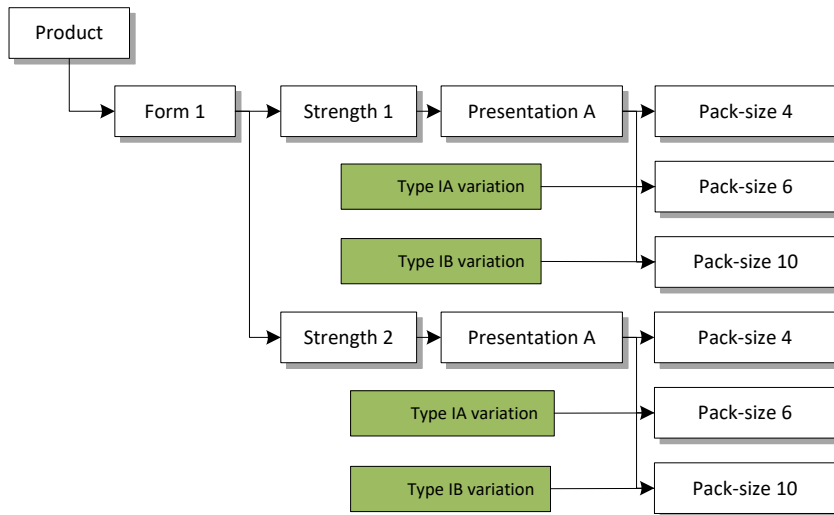
For a minor variation to a marketing authorisation, as defined in Article 2(2) of the Commission Regulation (EC) No 1234/2008 (type IA).



8 600 EURO

Basic fee

For a minor variation to a marketing authorisation, as defined in Article 2(5) of the Commission Regulation (EC) No 1234/2008 (type IB).



The addition of the 4 new pack sizes in the example above should be submitted as a grouping of 4 variations (2 x Type IA + 2 x Type IB).

Fee payable: 2 Type IA fees + 2 Type IB fees, i.e. 2 x 3 900 EURO + 2 x 8 600 EURO = 25 000 EURO

A.1.1.6. Examples of the determination of fees for renewals of marketing authorisation

It should be noted that the number of renewal fees charged for a medicinal product depends on the number of strengths associated with each pharmaceutical form determined as shown in section A.1.1.2. The number of presentations is not taken into consideration for the calculation of the renewal fee. The examples shown below do not represent an exhaustive list.

Scenario 1:

Full dossier application for a medicinal product having one pharmaceutical form with one strength and X presentations

Strengths associated with a pharmaceutical form:

- One strength associated with one pharmaceutical form



17 000 EURO

For renewal.

Basic fee

Scenario 2:

Full dossier application for a medicinal product having two pharmaceutical forms with two strengths and X presentations/strength associated with the first form and one strength and Y presentations associated with the second form.

Strengths associated with a pharmaceutical form:

- Two strengths associated with first pharmaceutical form
- One strength associated with second pharmaceutical form



2*17 000 EURO

For renewal.

Basic fee



+ 17 000 EURO

For renewal.

Additional fee

Scenario 3:

Full dossier application for an insulin product having two pharmaceutical forms with six strengths (consisting of two sets of one un-combined preparation and two combination preparations (having different proportions of insulin) with insulin amounts corresponding to A I.U. and B I.U.) and X presentations/ strength associated with the first form; and two strengths (of un-combined preparations with insulin amounts corresponding to A I.U. and B I.U.) and Y presentations/strength associated with the second form

Strengths associated with a pharmaceutical form:

- Six strengths associated with first pharmaceutical form
- Two strengths associated with second pharmaceutical form

**6*17 000 EURO**

For renewal.

Basic fee

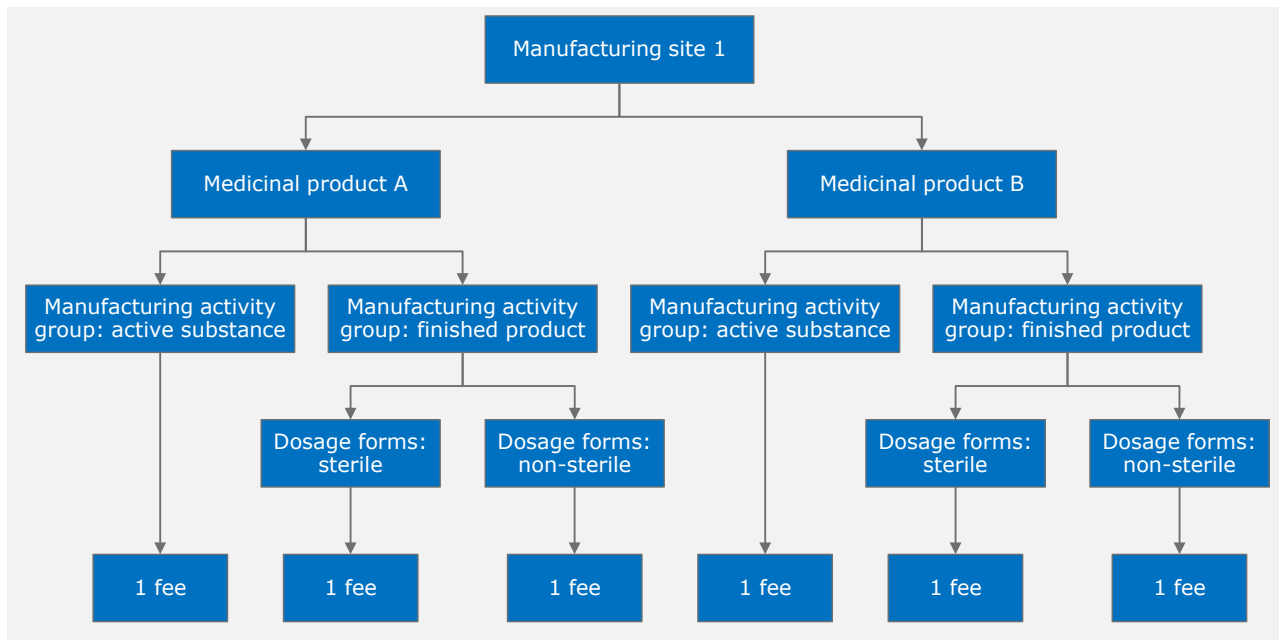
**+ 2*17 000 EURO**

For renewal.

Additional fee

A.1.1.7.1. Examples of the determination of fees for GMP inspections

In accordance with Annex IV to the “Rules for the implementation of Council Regulation (EC) No 297/95 on fees payable to the European Medicines Agency and other measures”, applicants are liable for more than one inspection fee on the basis of the following flowchart.



The examples shown below do not represent an exhaustive list. They apply to good manufacturing practice (GMP) inspections in relation to medicines for human use and to medicines for veterinary use.

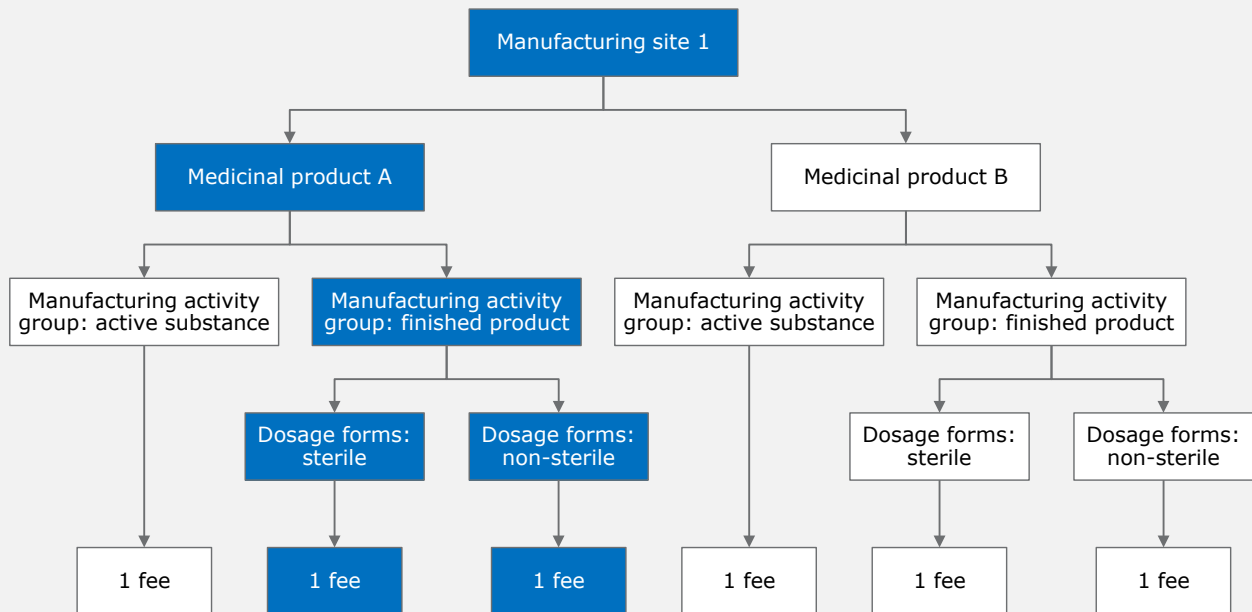
Scenario 1:

GMP inspection of manufacturing site 1 for one medicinal product A and involving two pharmaceutical forms: capsules (non-sterile) and solution for injection (sterile). The manufacturing activity for the two pharmaceutical forms is the same, i.e. manufacture of the finished product.

Fee payable: 2x basic fee (level I) 26 200 EURO = 52 400 EURO

Rationale: there are two types of dosages forms (sterile and non-sterile) and each one attracts a basic fee (Level I).

The applicable fees to be paid are represented by the blue boxes in the flowchart below.



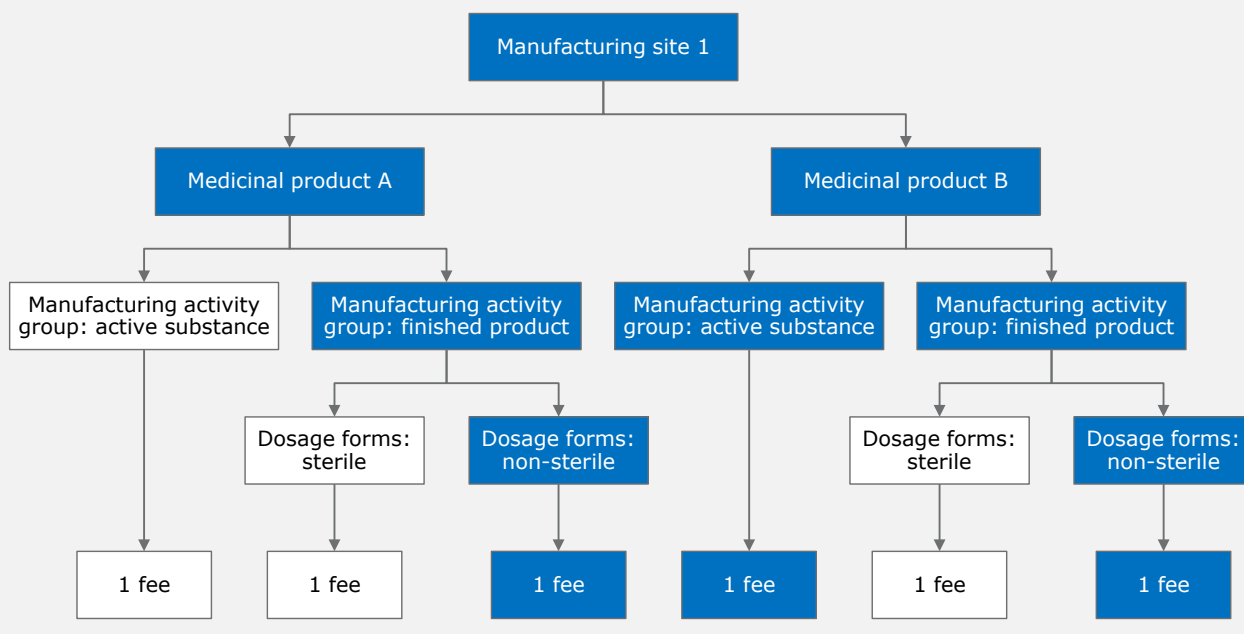
Scenario 2:

GMP inspection of manufacturing site 1 for two medicinal products (A and B). Product A involves only one pharmaceutical form (capsules) and one pharmaceutical activity (primary packaging). Product B also involves one pharmaceutical form (tablets) and four manufacturing activities (manufacture of the active substance, quality control of the active substance, manufacture of the finished product and primary packaging).

Fee payable: 3x basic fee (Level I) 26 200 EURO = 78 600 EURO; the applicable fees to be paid are represented by the blue boxes in the flowchart below.

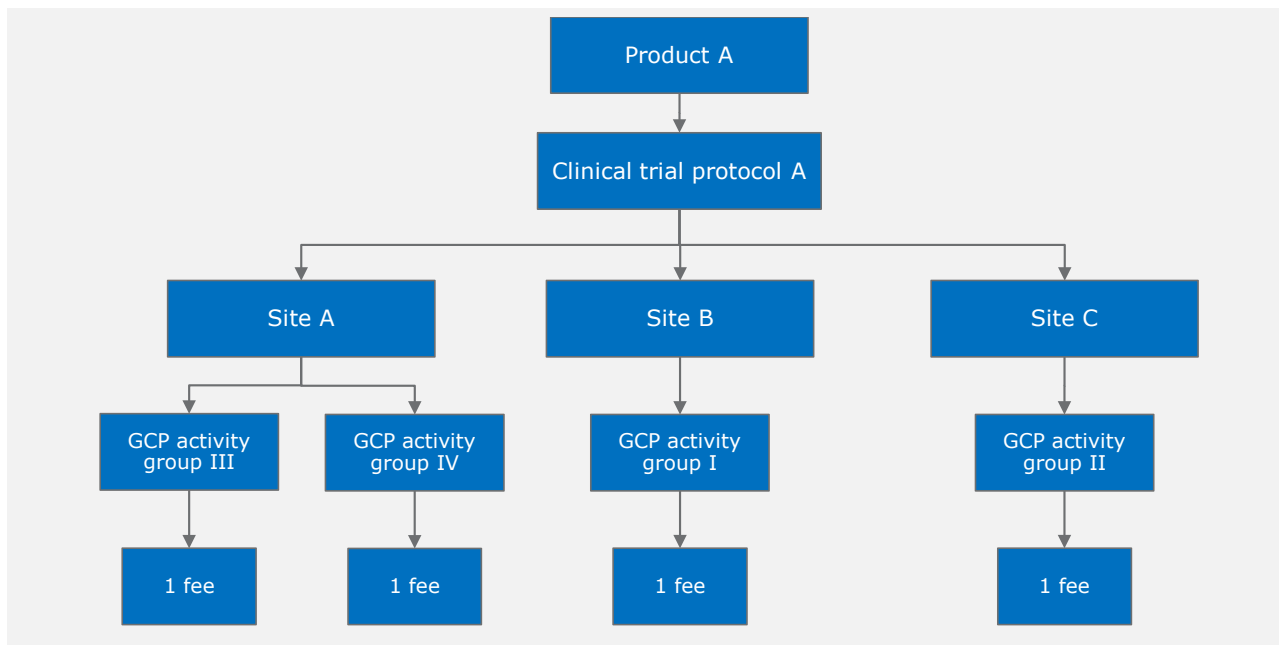
Rationale: Product A attracts only one fee because there is only one group of manufacturing activities (i.e. finished product) and one dosage form (non-sterile). Product B attracts two fees because there are manufacturing activities related to each group as follows:

- Group Active Substance: manufacture of the active substance and quality control of the active substance
- Group Finished Product: manufacture of the finished product and Primary Packaging



A.1.1.7.2. Examples of the determination of fees for GCP inspections

In accordance with Annex IV to the “Rules for the implementation of Council Regulation (EC) No 297/95 on fees payable to the European Medicines Agency and other measures”, applicants are liable for more than one inspection fee on the basis of the following flowchart.



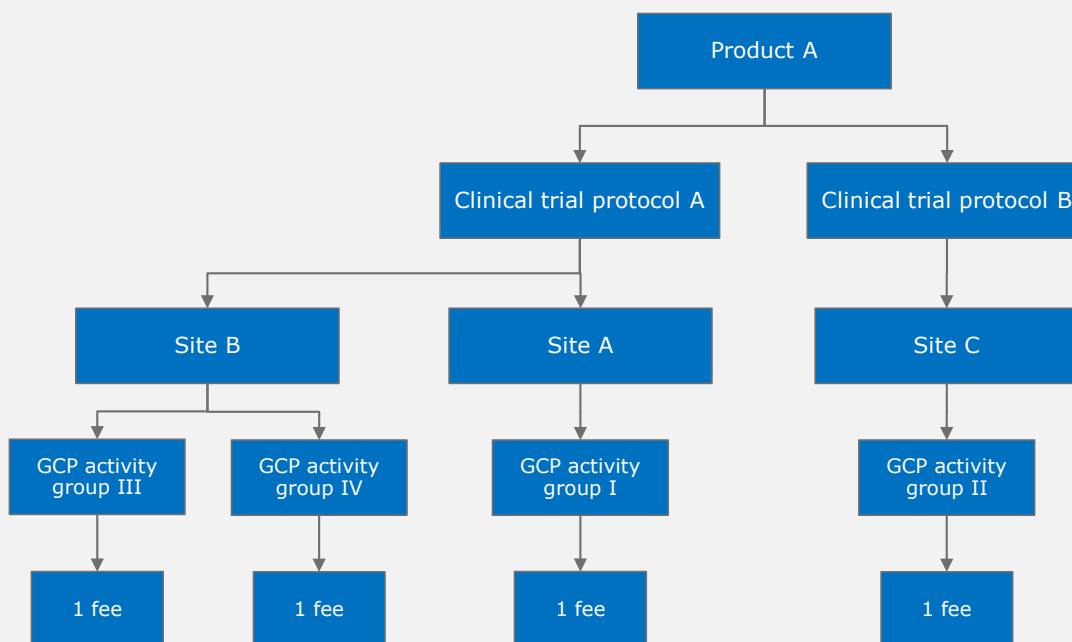
The examples shown below do not represent an exhaustive list. They apply to good clinical practice (GCP) inspections in relation to medicines for human use and to medicines for veterinary use.

Scenario 1:

GCP inspection request for Product A marketing authorisation application and the conduct of clinical trial protocol A at Site A (investigator site) for one clinical trial activity (Activity Group I) and at site B (sponsor site) for two clinical trial activities (Activity Group III and Activity Group IV). GCP inspection of clinical trial protocol B conducted at site C (Central laboratory) for one activity group (Activity group II);

Fee payable: 4x basic fee 26 200 EURO = 104 800 EURO

Rationale: there is one product A supported by two pivotal trials the conduct of which is inspected for clinical trial protocol A at Site A for one activity group (Activity Group I) and at site B for two activity Groups (Activity group III and Activity Group IV). The clinical trial protocol B is inspected at site C for one activity group (Activity Group II) and each activity group attracts a basic fee (Level I).

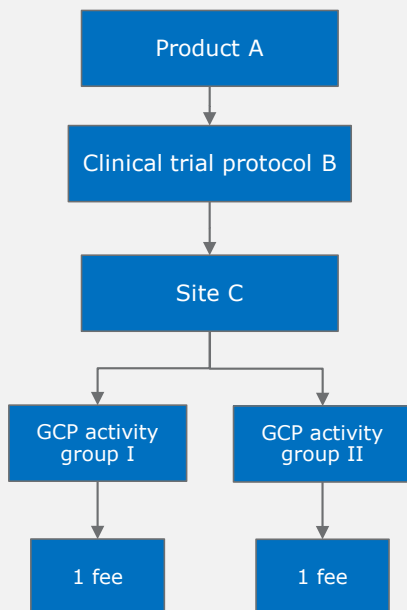


Scenario 2:

GCP inspection request for Product A marketing authorisation application and the conduct of clinical trial protocol B at Site C (i.e. CRO site including clinical and bioanalytical facility) for two clinical trial activities (Activity Group I) and (Activity Group II).

Fee payable: 2x basic fee 26 200 EURO = 52 400 EURO

Rationale: there is one product A supported by one pivotal trial the conduct of which is inspected for clinical trial protocol B at Site C for two activity groups (Activity Group I and Activity Group II) and each activity group attracts a basic fee (Level I).

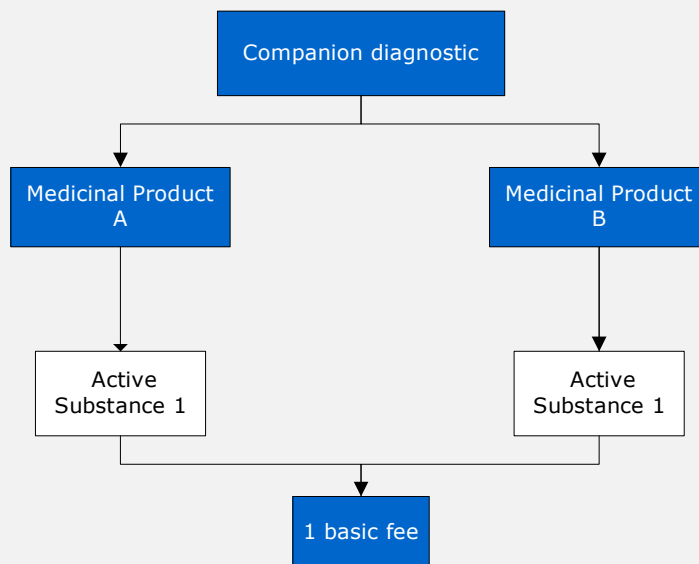


A.1.4.3 Examples of the determination of fees for applications for companion diagnostics devices

Scenario 1:

Initial request for a consultation on the suitability of a companion diagnostic for Product A containing active substance 1 and product B containing same active substance

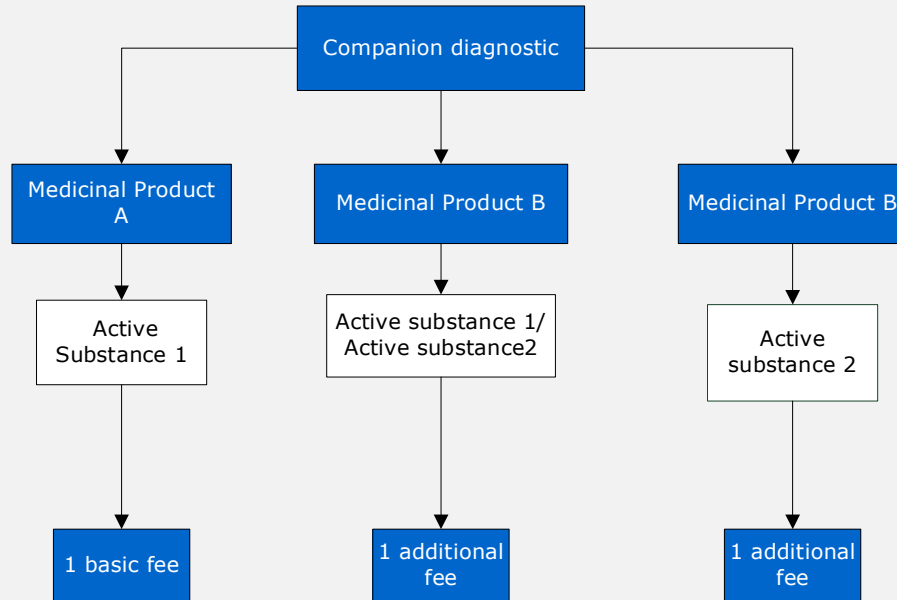
Fee payable: one basic fee, i.e., 51 800 EUR



Scenario 2:

Initial request for a consultation on the suitability of a companion diagnostic with medicinal product A containing active substance 1 and medicinal product B containing a combination of active substances 1 and 2 and medicinal product C containing active substance 2

Fee payable: one basic fee and one additional fee per active substance, i.e., 51 800 EUR + 8 600 EUR + 8 600 EUR = 69 000 EUR



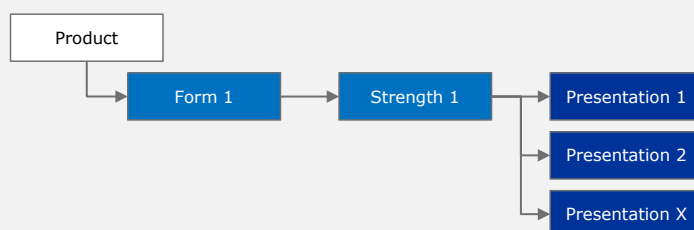
A.2. Veterinary medicinal products

A.2.1.2. Examples of the determination of fees for applications for marketing authorisation

It should be noted that the calculation of the total fee for a marketing authorisation application is driven by the pharmaceutical forms, the associated strengths and the associated presentations. The number of target species is irrelevant. The examples shown below do not represent an exhaustive list.

Scenario 1:

Full dossier application for a medicinal product having one pharmaceutical form with one strength and X presentations



173 000 EURO

Basic fee

Includes one pharmaceutical form and one associated strength and one presentation.



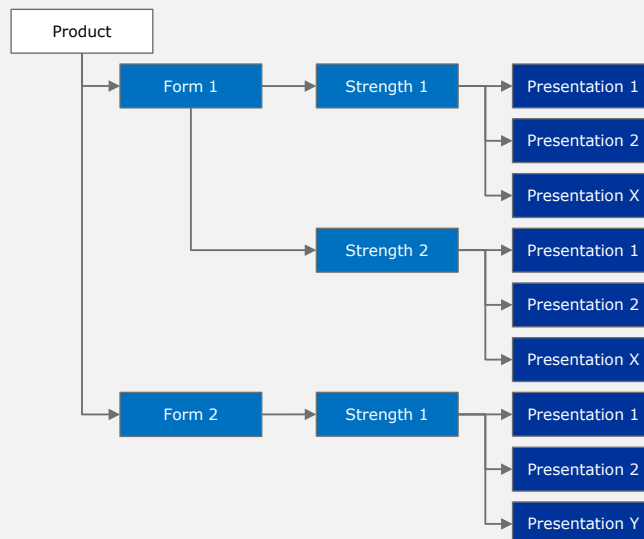
+ (X-1)*8 600 EURO

Additional fee

For additional presentations associated with the single strength.

Scenario 2:

Full dossier application for a medicinal product having two pharmaceutical forms with two strengths and X presentations/strength associated with the first form and one strength and Y presentations associated with the second form

**173 000 EURO**

Basic fee

Includes one pharmaceutical form and one associated strength and one presentation.

**+ (X-1)*8 600 EURO**

Additional fee

For additional presentations associated with the first form and strength.

**+ 17 000 EURO**

Additional fee

Second strength associated with the first form including one presentation.

**+ (X-1)*8 600 EURO**

Additional fee

For additional presentations associated with the first form and second strength.

**+ 17 000 EURO**

Additional fee

Second form including its associated strength and one presentation.

**+ (Y-1)*8 600 EURO**

Additional fee

For additional presentations associated with the second form and its strength.

A.2.1.4. Examples of the determination of fees for VRA level 1 of marketing authorisation


It should be noted that the basic fee for a VRA level 1 of a marketing authorisation is driven by the pharmaceutical form. The examples shown below do not represent an exhaustive list.


Scenario 1:


New pharmaceutical form with two strengths and X presentations/strength, for authorised or new route of administration (with submitted/cross-referenced clinical data)


VRA level 1 application:

- One pharmaceutical form, first strength and X presentations
- Second strength (of same new pharmaceutical form) and X presentations

 **43 300 EURO**
For VRA Level 1.
Basic fee

 **+ (X-1)*8 600 EURO**
For additional presentation fees.
Additional fee

 **+ 10 700 EURO**
For additional strength fee.
Additional fee


 **+ (X-1)*8 600 EURO**
For additional presentation fees.
Additional fee


Scenario 2:

New route of administration for authorised pharmaceutical form with two authorised strengths and X presentations/strength (with submitted/cross-referenced clinical data)

VRA level 1 application:

- Route of administration for authorised pharmaceutical form, first strength and X presentations
- Second strength (same new route of administration for same authorised pharmaceutical form) and X presentations

 **43 300 EURO**
For VRA Level 1.
Basic fee

 **+ (X-1)*8 600 EURO**
For additional presentation fees.
Additional fee



+ 10 700 EURO

Additional fee

For additional strength fee.



+ (X-1)*8 600 EURO

Additional fee

For additional presentation fees.

Scenario 3:

Two new strengths of same authorised pharmaceutical form and X presentations/strength (without submitted/cross-referenced clinical data)

VRA level 1 application:

- First new strength and X presentations
- Second new strength (of same authorised pharmaceutical form) and X presentations



39 100 EURO

Basic fee

For VRA Level 1.



+ (X-1)* 8 600 EURO

Additional fee

For additional presentation fees.



+ 10 700 EURO

Additional fee

For additional strength fee.



+ (X-1)* 8 600 EURO

Additional fee

For additional presentation fees.


Scenario 4:


One new strength of each of two authorised pharmaceutical forms and X presentations/strength (without submitted/cross-referenced clinical data)

THESE SHOULD BE SUBMITTED AS SEPARATE VRA LEVEL 1 APPLICATIONS:

VRA level 1 application 1:


- New strength (of first authorised pharmaceutical form) and X presentations


 **39 100 EURO**
Basic fee For VRA Level 1.

 **+ (X-1)*8 600 EURO**
Additional fee For additional presentation fees.

VRA level 1 application 2:


- New strength (of second authorised pharmaceutical form) and X presentations


 **39 100 EURO**
Basic fee For VRA Level 1.


 **+ (X-1)*8 600 EURO**
Additional fee For additional presentation fees.


VRA level 1 application 3:

- New pharmaceutical form for new target species, first strength and X presentations
- Second strength (of same new pharmaceutical form for same new target species) and X presentations

 **43 300 EURO**
Basic fee For VRA Level 1.


 **+ (X-1)*8 600 EURO**
Additional fee For additional presentation fees.


 **+ 10 700 EURO**
Additional fee For additional strength fee.

 **+ (X-1)*8 600 EURO**
Additional fee For additional presentation fees.

VRA level 1 application 4:


- New pharmaceutical form for a new target species, one strength and X presentations


 **43 300 EURO**
For VRA Level 1.
Basic fee


 **+ (X-1)*8 600 EURO**
For additional presentation fees.
Additional fee


VRA level 1 application 5:

- Second new target species of authorised pharmaceutical form, new strength and X presentations
- Second strength (for same new target species of authorised pharmaceutical form) and X presentations

 **43 300 EURO**
For VRA Level 1.
Basic fee

 **+ (X-1)*8 600 EURO**
For additional presentation fees.
Additional fee

 **+ 10 700 EURO**
For additional strength fee.
Additional fee

 **+ (X-1)*8 600 EURO**
For additional presentation fees.
Additional fee

A.3. Administrative fees

A.3.2. Examples of the determination of fees for certificates of medicinal product

In accordance with Annex III of the "Rules for the implementation of Council Regulation (EC) No 297/95 on fees payable to the European Medicines Agency and other measures", requesters are liable for more than one fee depending on the number of medicinal products, importing countries, languages and annexes.


The examples shown below do not represent an exhaustive list.


Scenario 1:


One request for certificates for one medicinal product, as follows:

- Addressed to country 1: 5 certificates with annex 1
- Addressed to country 2: 10 certificates with annex 1


Fee payable using the standard procedure: **720 EURO**


 **360 EURO**
Basic fee For the first set of 5 certificates for country 1.


 **+ 180 EURO**
Additional fee For the second set of maximum 6 out of 10 certificates for country 2.

 **+ 180 EURO**
Additional fee For the third set of 4 out of 10 certificates for country 2.

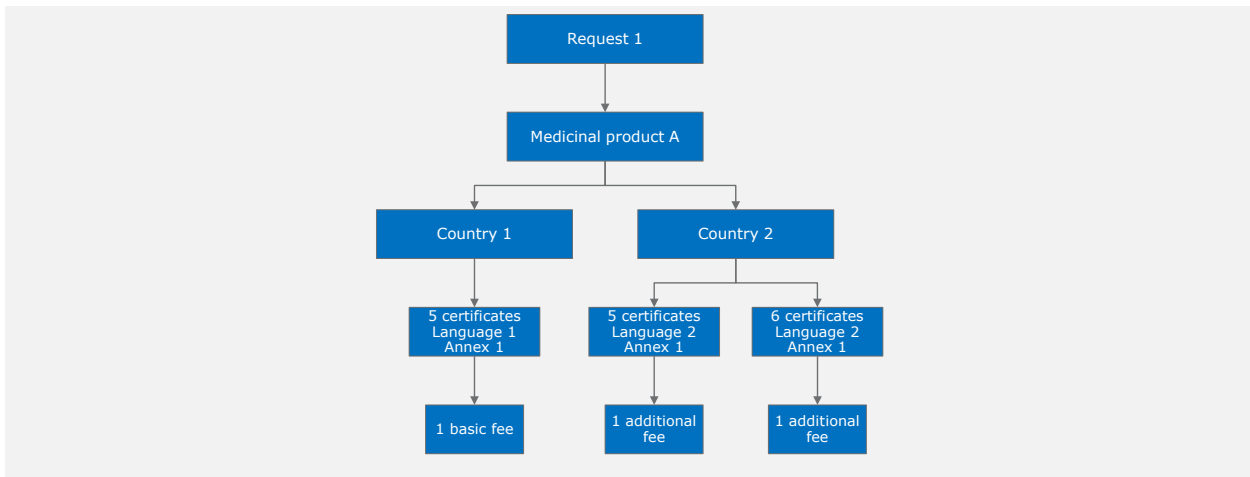
Fee payable using the urgent procedure: **2 160 EURO**

 **1 080 EURO**
Basic fee For the first set of 5 certificates for country 1.

 **+ 540 EURO**
Additional fee For the second set of maximum 6 out of 10 certificates for country 2.

 **+ 540 EURO**
Additional fee For the third set of 4 out of 10 certificates for country 2.

The applicable fees to be paid using the standard procedure are represented in the flowchart below.





Scenario 2:


One request for certificates for two medicinal products, as follows:


- Medicinal product A:
 - Addressed to country 1: 5 certificates with annex 1
 - Addressed to country 2: 6 certificates with Annex 1, 9 certificates with annexes 1 and 2
- Medicinal product B:
 - Addressed to country 1: 15 certificates with annex 1
 - Addressed to country 2: 6 certificates with annex 1


Fee payable using the standard procedure: **1 620 EURO** basic fee + (additional fee x 7)


 **360 EURO**
Basic fee
For the first set of 5 certificates for medicinal product A for country 1.

 **+ 180 EURO**
Additional fee
For the second set of 6 certificates for medicinal product A for country 2.

 **+ 180 EURO**
Additional fee
For the third set of maximum 6 out of 9 certificates for medicinal product A for country 2.

 **+ 180 EURO**
Additional fee
For the fourth set of 3 out of 9 certificates for medicinal product A for country 2.

 **+ 180 EURO**
Additional fee
For the fifth set of maximum 6 out of 15 certificates for medicinal product B for country 1.

 **+ 180 EURO**
For the sixth set of maximum 6 out of 15 certificates for medicinal product B for

Additional fee country 1.



+ 180 EURO

Additional fee

For the seventh set of 3 out of 15 certificates for medicinal product B for country 1.



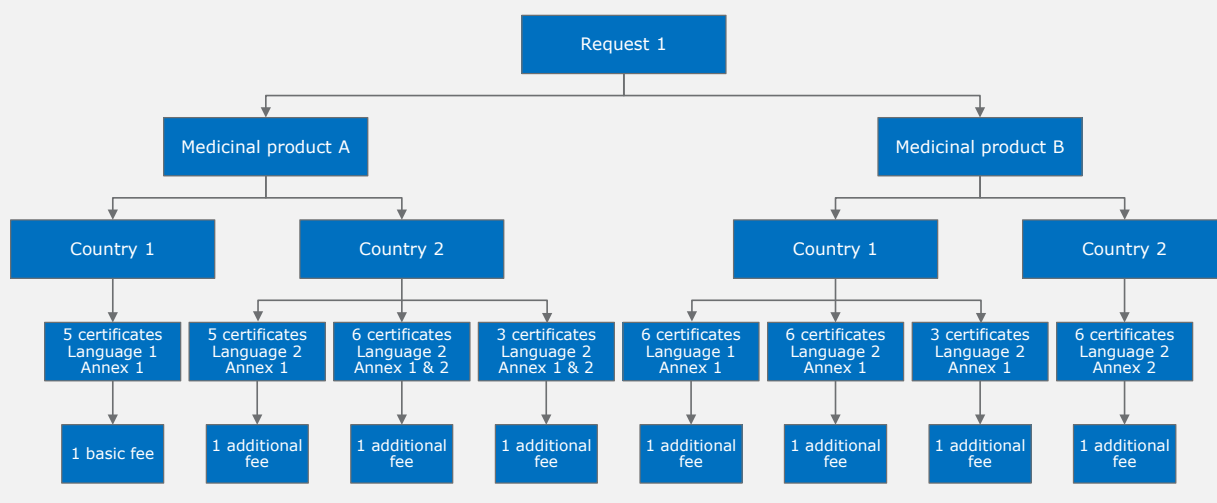
+ 180 EURO

Additional fee

For the eighth set of 6 certificates for medicinal product B for country 2.

Total fee payable using the urgent procedure: applying the same rationale as for the standard procedure, one basic fee and seven additional fees are payable: = 1 080 EURO + (540 EURO* 7)
= 4 860 EURO

The applicable fees to be paid using the standard procedure are represented in the flowchart below



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